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(54) Title: PRODUCTION OF POLYUNSATURATED FATTY ACIDS BY EXPRESSION OF POLYKETIDE-LIKE SYNTHESIS GENES IN PLANTS					
(57) Abstract <p>The present invention relates to compositions and methods for preparing polyunsaturated long chain fatty acids in plants, plant parts and plant cells, such as leaves, roots, fruits and seeds. Nucleic acid sequences and constructs encoding PKS-like genes required for the poly-unsaturated long chain fatty acid production, including the genes responsible for eicosapentenoic acid production of <i>Shewanella putrefaciens</i> and novel genes associated with the production of docosahexenoic acid in <i>Vibrio marinus</i> are used to generate transgenic plants, plant parts and cells which contain and express one or more transgenes encoding one or more of the PKS-like genes associated with such long chain polyunsaturated fatty acid production. Expression of the PKS-like genes in the plant system permits the large scale production of polyunsaturated long chain fatty acids such as eicosapentenoic acid and docosahexenoic acid for modification of the fatty acid profile of plants, plant parts and tissues. Manipulation of the fatty acid profiles allows for the production of commercial quantities of novel plant oils and products.</p>					

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PRODUCTION OF POLYUNSATURATED FATTY ACIDS BY EXPRESSION OF POLYKETIDE-LIKE SYNTHESIS GENES IN PLANTS

INTRODUCTION

5 Field of the Invention

This invention relates to modulating levels of enzymes and/or enzyme components capable of modifying long chain poly-unsaturated fatty acids (PUFAs) in a host cell, and constructs and methods for producing PUFAs in a host cell. The invention is exemplified by production of eicosapentenoic acid (EPA) using genes derived from *Shewanella* 10 *putrefaciens* and *Vibrio marinus*.

Background

Two main families of poly-unsaturated fatty acids (PUFAs) are the $\omega 3$ fatty acids, exemplified by eicosapentenoic acid, and the $\omega 6$ fatty acids, exemplified by arachidonic acid. PUFAs are important components of the plasma membrane of the cell, where they can be found in such forms as phospholipids, and also can be found in triglycerides. PUFAs also serve as precursors to other molecules of importance in human beings and animals, including the prostacyclins, leukotrienes and prostaglandins. Long chain PUFAs of importance include docosahexenoic acid (DHA) and eicosapentenoic acid (EPA), 20 which are found primarily in different types of fish oil, gamma-linolenic acid (GLA), which is found in the seeds of a number of plants, including evening primrose (*Oenothera biennis*), borage (*Borago officinalis*) and black currants (*Ribes nigrum*), stearidonic acid (SDA), which is found in marine oils and plant seeds, and arachidonic acid (ARA), which along with GLA is found in filamentous fungi. ARA can be purified from animal tissues 25 including liver and adrenal gland. Several genera of marine bacteria are known which synthesize either EPA or DHA. DHA is present in human milk along with ARA.

PUFAs are necessary for proper development, particularly in the developing infant brain, and for tissue formation and repair. As an example, DHA, is an important constituent of many human cell membranes, in particular nervous cells (gray matter), 30 muscle cells, and spermatozoa and believed to affect the development of brain functions in general and to be essential for the development of eyesight. EPA and DHA have a number of nutritional and pharmacological uses. As an example adults affected by diabetes (especially non insulin-dependent) show deficiencies and imbalances in their

levels of DHA which are believed to contribute to later coronary conditions. Therefore a diet balanced in DHA may be beneficial to diabetics.

For DHA, a number of sources exist for commercial production including a variety of marine organisms, oils obtained from cold water marine fish, and egg yolk fractions. The purification of DHA from fish sources is relatively expensive due to technical difficulties, making DHA expensive and in short supply. In algae such as *Amphidinium* and *Schyzochytrium* and marine fungi such as *Thraustochytrium* DHA may represent up to 48% of the fatty acid content of the cell. A few bacteria also are reported to produce DHA. These are generally deep sea bacteria such as *Vibrio marinus*. For ARA, microorganisms including the genera *Mortierella*, *Entomophthora*, *Phytium* and *Porphyridium* can be used for commercial production. Commercial sources of SDA include the genera *Trichodesma* and *Echium*. Commercial sources of GLA include evening primrose, black currants and borage. However, there are several disadvantages associated with commercial production of PUFAs from natural sources. Natural sources of PUFA, such as animals and plants, tend to have highly heterogeneous oil compositions. The oils obtained from these sources can require extensive purification to separate out one or more desired PUFA or to produce an oil which is enriched in one or more desired PUFA.

Natural sources also are subject to uncontrollable fluctuations in availability. Fish stocks may undergo natural variation or may be depleted by overfishing. Animal oils, and particularly fish oils, can accumulate environmental pollutants. Weather and disease can cause fluctuation in yields from both fish and plant sources. Cropland available for production of alternate oil-producing crops is subject to competition from the steady expansion of human populations and the associated increased need for food production on the remaining arable land. Crops which do produce PUFAs, such as borage, have not been adapted to commercial growth and may not perform well in monoculture. Growth of such crops is thus not economically competitive where more profitable and better established crops can be grown. Large -scale fermentation of organisms such as *Shewanella* also is expensive. Natural animal tissues contain low amounts of ARA and are difficult to process. Microorganisms such as *Porphyridium* and *Shewanella* are difficult to cultivate on a commercial scale.

Dietary supplements and pharmaceutical formulations containing PUFAs can retain the disadvantages of the PUFA source. Supplements such as fish oil capsules can

contain low levels of the particular desired component and thus require large dosages. High dosages result in ingestion of high levels of undesired components, including contaminants. Care must be taken in providing fatty acid supplements, as overaddition may result in suppression of endogenous biosynthetic pathways and lead to competition with other necessary fatty acids in various lipid fractions *in vivo*, leading to undesirable results. For example, Eskimos having a diet high in $\omega 3$ fatty acids have an increased tendency to bleed (U.S. Pat. No. 4,874,603). Fish oils have unpleasant tastes and odors, which may be impossible to economically separate from the desired product, such as a food supplement. Unpleasant tastes and odors of the supplements can make such regimens involving the supplement undesirable and may inhibit compliance by the patient.

A number of enzymes have been identified as being involved in PUFA biosynthesis. Linoleic acid (LA, 18:2 Δ 9, 12) is produced from oleic acid (18:1 Δ 9) by a $\Delta 12$ -desaturase. GLA (18:3 Δ 6, 9, 12) is produced from linoleic acid (LA, 18:2 Δ 9, 12) by a $\Delta 6$ -desaturase. ARA (20:4 Δ 5, 8, 11, 14) is produced from DGLA (20:3 Δ 8, 11, 14), catalyzed by a $\Delta 5$ -desaturase. Eicosapentenoic acid (EPA) is a 20 carbon, omega 3 fatty acid containing 5 double bonds (Δ 5, 8, 11, 14, 17), all in the *cis* configuration. EPA, and the related DHA (Δ 4, 7, 10, 13, 16, 19, C22:6) are produced from oleic acid by a series of elongation and desaturation reactions. Additionally, an elongase (or elongases) is required to extend the 18 carbon PUFAs out to 20 and 22 carbon chain lengths.

However, animals cannot convert oleic acid (18:1 Δ 9) into linoleic acid (18:2 Δ 9, 12). Likewise, μ -linolenic acid (ALA, 18:3 Δ 9, 12, 15) cannot be synthesized by mammals. Other eukaryotes, including fungi and plants, have enzymes which desaturate at positions $\Delta 12$ and $\Delta 15$. The major poly-unsaturated fatty acids of animals therefore are either derived from diet and/or from desaturation and elongation of linoleic acid (18:2 Δ 9, 12) or μ -linolenic acid (18:3 Δ 9, 12, 15).

Poly-unsaturated fatty acids are considered to be useful for nutritional, pharmaceutical, industrial, and other purposes. An expansive supply of poly-unsaturated fatty acids from natural sources and from chemical synthesis are not sufficient for commercial needs. Because a number of separate desaturase and elongase enzymes are required for fatty acid synthesis from linoleic acid (LA, 18:2 Δ 9, 12), common in most plant species, to the more saturated and longer chain PUFAs, engineering plant host cells for the expression of EPA and DHA may require expression of five or six separate

enzyme activities to achieve expression, at least for EPA and DHA, and for production of quantities of such PUFAs additional engineering efforts may be required, for instance the down regulation of enzymes competing for substrate, engineering of higher enzyme activities such as by mutagenesis or targeting of enzymes to plastid organelles. Therefore 5 it is of interest to obtain genetic material involved in PUFA biosynthesis from species that naturally produce these fatty acids and to express the isolated material alone or in combination in a heterologous system which can be manipulated to allow production of commercial quantities of PUFAs.

10 Relevant Literature

Several genera of marine bacteria have been identified which synthesize either EPA or DHA (DeLong and Yayanos, *Applied and Environmental Microbiology* (1986) 51: 730-737). Researchers of the Sagami Chemical Research Institute have reported EPA production in *E. coli* which have been transformed with a gene cluster from the marine 15 bacterium, *Shewanella putrefaciens*. A minimum of 5 open reading frames (ORFs) are required for fatty acid synthesis of EPA in *E. coli*. To date, extensive characterization of the functions of the proteins encoded by these genes has not been reported (Yazawa (1996) *Lipids* 31, S-297; WO 93/23545; WO 96/21735).

The protein sequence of open reading frame (ORF) 3 as published by Yazawa, 20 USPN 5,683,898 is not a functional protein. Yazawa defines the protein as initiating at the methionine codon at nucleotides 9016-9014 of the *Shewanella* PKS-like cluster (Genbank accession U73935) and ending at the stop codon at nucleotides 8185-8183 of the *Shewanella* PKS-like cluster. However, when this ORF is expressed under control of a heterologous promoter in an *E. coli* strain containing the entire PKS-like cluster except 25 ORF 3, the recombinant cells do not produce EPA.

Polyketides are secondary metabolites the synthesis of which involves a set of enzymatic reactions analogous to those of fatty acid synthesis (see reviews: Hopwood and Sherman, *Annu. Rev. Genet.* (1990) 24: 37-66, and Katz and Donadio, *in Annual Review of Microbiology* (1993) 47: 875-912). It has been proposed to use polyketide 30 synthases to produce novel antibiotics (Hutchinson and Fujii, *Annual Review of Microbiology* (1995) 49:201-238).

SUMMARY OF THE INVENTION

Novel compositions and methods are provided for preparation of long chain poly-
unsaturated fatty acids (PUFAs) using polyketide-like synthesis (PKS-like) genes in
5 plants and plant cells. In contrast to the known and proposed methods for production of
PUFAs by means of fatty acid synthesis genes, by the invention constructs and methods
are provided for producing PUFAs by utilizing genes of a PKS-like system. The methods
involve growing a host cell of interest transformed with an expression cassette functional
in the host cell, the expression cassette comprising a transcriptional and translational
10 initiation regulatory region, joined in reading frame 5' to a DNA sequence to a gene or
component of a PKS-like system capable of modulating the production of PUFAs (PKS-
like gene). An alteration in the PUFA profile of host cells is achieved by expression
following introduction of a complete PKS-like system responsible for a PUFA
biosynthesis into host cells. The invention finds use for example in the large scale
15 production of DHA and EPA and for modification of the fatty acid profile of host cells
and edible plant tissues and/or plant parts.

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 provides designations for the ORFs of the EPA gene cluster of
20 *Shewanella*. Figure 1A shows the organization of the genes; those ORFs essential for
EPA production in *E. coli* are numbered. Figure 1B shows the designations given to
subclones.

Figure 2 provides the *Shewanella* PKS-like domain structure, motifs and 'Blast'
matches of ORF 6 (Figure 2A), ORF 7 (Figure 2B), ORF 8 (Figure 2C), ORF 9
25 (Figure 2D) and ORF 3 (Figure 2E). Figure 2F shows the structure of the region of the
Anabeana chromosome that is related to domains present in *Shewanella* EPA ORFs.

Figure 3 shows results for pantethenylation - ORF 3 in *E. coli* strain SJ16.

Figure 4 is the sequence for the PKS-like cluster found in *Shewanella*, containing
30 ORFs 3, 4, 5, 6, 7, 8 and 9. The start and last codons for each ORF are as follows:
ORF3 (published-inactive): 9016, 8186; ORF3 (active in EPA synthesis): 9157, 8186;
ORF 6: 13906, 22173; ORF 7: 22203, 24515; ORF 8: 24518, 30529; ORF 9: 30730,
32358.

Figure 5 shows the sequence for the PKS-like cluster in an approximately 40 kb DNA fragment of *Vibrio marinus*, containing ORFs 6, 7, 8 and 9. The start and last condons for each ORF are as follows: ORF 6: 17394, 25352; ORF 7: 25509, 28160; ORF 8: 28209, 34265; ORF 9: 34454, 36118.

5 Figure 6 shows the sequence for an approximately 19 kb portion of the PKS-like cluster of Figure 5 which contains the ORFs 6, 7, 8 and 9. The start and last condons for each ORF are as follows: ORF 6: 411, 8369; ORF 7: 8526, 11177; ORF 8: 11226, 17282; ORF 9: 17471, 19135.

10 Figure 7 shows a comparison of the PKS-like gene clusters of *Shewanella putrefaciens* and *Vibrio marinus*; Figure 7B is the *Vibrio marinus* operon sequence.

Figure 8 is an expanded view of the PKS-like gene cluster portion of *Vibrio marinus* shown in Figure 7B showing that ORFs 6, 7 and 8 are in reading frame 2, while ORF 9 is in reading frame 3.

15 Figure 9 demonstrates sequence homology of ORF 6 of *Shewanella putrefaciens* and *Vibrio marinus*. The *Shewanella* ORF 6 is depicted on the vertical axis, and the *Vibrio* ORF 6 is depicted on the horizontal axis. Lines indicate regions of the proteins that have a 60% identity. The repeated lines in the middle correspond to the multiple ACP domains found in ORF 6.

20 Figure 10 demonstrates sequence homology of ORF 7 of *Shewanella putrefaciens* and *Vibrio marinus*. The *Shewanella* ORF 7 is depicted on the vertical axis, and the *Vibrio* ORF 7 is depicted on the horizontal axis. Lines indicate regions of the proteins that have a 60% identity.

25 Figure 11 demonstrates sequence homology of ORF 8 of *Shewanella putrefaciens* and *Vibrio marinus*. The *Shewanella* ORF 8 is depicted on the vertical axis, and the *Vibrio*. ORF 8 is depicted on the horizontal axis. Lines indicate regions of the proteins that have a 60% identity.

30 Figure 12 demonstrates sequence homology of ORF 9 of *Shewanella putrefaciens* and *Vibrio marinus*. The *Shewanella* ORF 9 is depicted on the vertical axis, and the *Vibrio* ORF 9 is depicted on the horizontal axis. Lines indicate regions of the proteins that have a 60% identity.

Figure 13 is a depiction of various complementation experiments, and resulting PUFA production. On the right, is shown the longest PUFA made in the *E. coli* strain

containing the *Vibrio* and *Shewanella* genes depicted on the left. The hollow boxes indicate ORFs from *Shewanella*. The solid boxes indicate ORFs from *Vibrio*.

Figure 14 is a chromatogram showing fatty acid production from complementation of pEPAD8 from *Shewanella* (deletion ORF 8) with ORF 8 from *Shewanella*, in *E. coli* Fad E-. The chromatogram presents an EPA (20:5) peak.

Figure 15 is a chromatogram showing fatty acid production from complementation of pEPAD8 from *Shewanella* (deletion ORF 8) with ORF 8 from *Vibrio marinus*, in *E. coli* Fad E-. The chromatograph presents EPA (20:5) and DHA (22:6) peaks.

Figure 16 is a table of PUFA values from the ORF 8 complementation experiment, the chromatogram of which is shown in Figure 15.

Figure 17 is a plasmid map showing the elements of pCGN7770.

Figure 18 is a plasmid map showing the elements of pCGN8535.

Figure 19 is a plasmid map showing the elements of pCGN8537.

Figure 20 is a plasmid map showing the elements of pCGN8525.

Figure 21 is a comparison of the *Shewanella* ORFs as defined by Yazawa and those disclosed in Figure 4. When a protein starting at the leucine (TTG) codon at nucleotides 9157-9155 and ending at the stop codon at nucleotides 8185-8183 is expressed under control of a heterologous promoter in an *E. coli* strain containing the entire PKS-like cluster except ORF 3, the recombinant cells do produce EPA. Thus, the published protein sequence is likely to be wrong, and the coding sequence for the protein may start at the TTG codon at nucleotides 9157-9155 or the TTG codon at nucleotides 9172-9170. This information is critical to the expression of a functional PKS-like cluster heterologous system.

Figure 22 is a plasmid map showing the elements of pCGN8560.

Figure 23 is plasmid map showing the elements of pCGN8556.

Figure 24 shows the translated DNA sequence upstream of the published ORF 3. The ATG start codon at position 9016 is the start codon for the protein described by Yazawa *et al* (1996) *supra*. The other arrows depict TTG or ATT codons that can also serve as start codons in bacteria. When ORF 3 is started from the published ATG codon at 9016, the protein is not functional in making EPA. When ORF 3 is initiated at the TTG codon at position 9157, the protein is capable of facilitating EPA synthesis.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

In accordance with the subject invention, novel DNA sequences, DNA constructs and methods are provided, which include some or all of the polyketide-like synthesis (PKS-like) pathway genes from *Shewanella*, *Vibrio* or other microorganisms, for 5 modifying the poly-unsaturated long chain fatty acid content of host cells, particularly host plant cells. The present invention demonstrates that EPA synthesis genes in *Shewanella putrefaciens* constitute a polyketide-like synthesis pathway. Functions are ascribed to the *Shewanella* and *Vibrio* genes and methods are provided for the production 10 of EPA and DHA in host cells. The method includes the step of transforming cells with an expression cassette comprising a DNA encoding a polypeptide capable of increasing the amount of one or more PUFA in the host cell. Desirably, integration constructs are prepared which provide for integration of the expression cassette into the genome of a 15 host cell. Host cells are manipulated to express a sense or antisense DNA encoding a polypeptide(s) that has PKS-like gene activity. By "PKS-like gene" is intended a polypeptide which is responsible for any one or more of the functions of a PKS-like activity of interest. By "polypeptide" is meant any chain of amino acids, regardless of length or post-translational modification, for example, glycosylation or phosphorylation. Depending upon the nature of the host cell, the substrate(s) for the expressed enzyme may 20 be produced by the host cell or may be exogenously supplied. Of particular interest is the selective control of PUFA production in plant tissues and/or plant parts such as leaves, roots, fruits and seeds. The invention can be used to synthesize EPA, DHA, and other related PUFA in host cells.

There are many advantages to transgenic production of PUFA. As an example, in 25 transgenic *E. coli* as in *Shewanella*, EPA accumulates in the phospholipid fraction, specifically in the *sn*-2 position. It may be possible to produce a structured lipid in a desired host cell which differs substantially from that produced in either *Shewanella* or *E. coli*. Additionally transgenic production of PUFA in particular host cells offers several 30 advantages over purification from natural sources such as fish or plants. In transgenic plants, by utilizing a PKS-like system, fatty acid synthesis of PUFA is achieved in the cytoplasm by a system which produces the PUFA through *de novo* production of the fatty acids utilizing malonyl Co-A and acetyl Co-A as substrates. In this fashion, potential problems, such as those associated with substrate competition and diversion of normal products of fatty acid synthesis in a host to PUFA production, are avoided.

Production of fatty acids from recombinant plants provides the ability to alter the naturally occurring plant fatty acid profile by providing new synthetic pathways in the host or by suppressing undesired pathways, thereby increasing levels of desired PUFAs, or conjugated forms thereof, and decreasing levels of undesired PUFAs. Production of fatty acids in transgenic plants also offers the advantage that expression of PKS-like genes in particular tissues and/or plant parts means that greatly increased levels of desired PUFAs in those tissues and/or parts can be achieved, making recovery from those tissues more economical. Expression in a plant tissue and/or plant part presents certain efficiencies, particularly where the tissue or part is one which is easily harvested, such as seed, leaves, fruits, flowers, roots, etc. For example, the desired PUFAs can be expressed in seed; methods of isolating seed oils are well established. In addition to providing a source for purification of desired PUFAs, seed oil components can be manipulated through expression of PKS-like genes, either alone or in combination with other genes such as elongases, to provide seed oils having a particular PUFA profile in concentrated form. The concentrated seed oils then can be added to animal milks and/or synthetic or semisynthetic milks to serve as infant formulas where human nursing is impossible or undesired, or in cases of malnourishment or disease in both adults and infants.

Transgenic microbial production of fatty acids offers the advantages that many microbes are known with greatly simplified oil compositions as compared with those of higher organisms, making purification of desired components easier. Microbial production is not subject to fluctuations caused by external variables such as weather and food supply. Microbially produced oil is substantially free of contamination by environmental pollutants. Additionally, microbes can provide PUFAs in particular forms which may have specific uses. For example, *Spirulina* can provide PUFAs predominantly at the first and third positions of triglycerides; digestion by pancreatic lipases preferentially releases fatty acids from these positions. Following human or animal ingestion of triglycerides derived from *Spirulina*, these PUFAs are released by pancreatic lipases as free fatty acids and thus are directly available, for example, for infant brain development. Additionally, microbial oil production can be manipulated by controlling culture conditions, notably by providing particular substrates for microbially expressed enzymes, or by addition of compounds which suppress undesired biochemical pathways. In addition to these advantages, production of fatty acids from recombinant microbes provides the ability to alter the naturally occurring microbial fatty acid profile by

providing new synthetic pathways in the host or by suppressing undesired pathways, thereby increasing levels of desired PUFAs, or conjugated forms thereof, and decreasing levels of undesired PUFAs.

Production of fatty acids in animals also presents several advantages. Expression 5 of desaturase genes in animals can produce greatly increased levels of desired PUFAs in animal tissues, making recovery from those tissues more economical. For example, where the desired PUFAs are expressed in the breast milk of animals, methods of isolating PUFAs from animal milk are well established. In addition to providing a source for purification of desired PUFAs, animal breast milk can be manipulated through 10 expression of desaturase genes, either alone or in combination with other human genes, to provide animal milks with a PUFA composition substantially similar to human breast milk during the different stages of infant development. Humanized animal milks could serve as infant formulas where human nursing is impossible or undesired, or in the cases of malnourishment or disease.

15 DNAs encoding desired PKS-like genes can be identified in a variety of ways. In one method, a source of a desired PKS-like gene, for example genomic libraries from a *Shewanella* or *Vibrio* spp., is screened with detectable enzymatically- or chemically-synthesized probes. Sources of ORFs having PKS-like genes are those organisms which produce a desired PUFA, including DHA-producing or EPA-producing deep sea bacteria 20 growing preferentially under high pressure or at relatively low temperature.

Microorganisms such as *Shewanella* which produce EPA or DHA also can be used as a source of PKS-like genes. The probes can be made from DNA, RNA, or non-naturally occurring nucleotides, or mixtures thereof. Probes can be enzymatically synthesized from 25 DNAs of known PKS-like genes for normal or reduced-stringency hybridization methods. For discussions of nucleic acid probe design and annealing conditions, see, for example, Sambrook *et al*, *Molecular Cloning: A Laboratory Manual* (2nd ed.), Vols. 1-3, *Cold Spring Harbor Laboratory*, (1989) or *Current Protocols in Molecular Biology*, F. Ausubel *et al*, ed., Greene Publishing and Wiley-Interscience, New York (1987), each of 30 which is incorporated herein by reference. Techniques for manipulation of nucleic acids encoding PUFA enzymes such as subcloning nucleic acid sequences encoding polypeptides into expression vectors, labelling probes, DNA hybridization, and the like are described generally in Sambrook, *supra*.

Oligonucleotide probes also can be used to screen sources and can be based on sequences of known PKS-like genes, including sequences conserved among known PKS-like genes, or on peptide sequences obtained from a desired purified protein.

5 Oligonucleotide probes based on amino acid sequences can be degenerate to encompass the degeneracy of the genetic code, or can be biased in favor of the preferred codons of the source organism. Alternatively, a desired protein can be entirely sequenced and total synthesis of a DNA encoding that polypeptide performed.

Once the desired DNA has been isolated, it can be sequenced by known methods. It is recognized in the art that such methods are subject to errors, such that multiple 10 sequencing of the same region is routine and is still expected to lead to measurable rates of mistakes in the resulting deduced sequence, particularly in regions having repeated domains, extensive secondary structure, or unusual base compositions, such as regions with high GC base content. When discrepancies arise, resequencing can be done and can employ special methods. Special methods can include altering sequencing conditions by 15 using: different temperatures; different enzymes; proteins which alter the ability of oligonucleotides to form higher order structures; altered nucleotides such as ITP or methylated dGTP; different gel compositions, for example adding formamide; different primers or primers located at different distances from the problem region; or different templates such as single stranded DNAs. Sequencing of mRNA can also be employed.

20 For the most part, some or all of the coding sequences for the polypeptides having PKS-like gene activity are from a natural source. In some situations, however, it is desirable to modify all or a portion of the codons, for example, to enhance expression, by employing host preferred codons. Host preferred codons can be determined from the 25 codons of highest frequency in the proteins expressed in the largest amount in a particular host species of interest. Thus, the coding sequence for a polypeptide having PKS-like gene activity can be synthesized in whole or in part. All or portions of the DNA also can be synthesized to remove any destabilizing sequences or regions of secondary structure which would be present in the transcribed mRNA. All or portions of the DNA also can be synthesized to alter the base composition to one more preferable to the desired host 30 cell. Methods for synthesizing sequences and bringing sequences together are well established in the literature. *In vitro* mutagenesis and selection, site-directed mutagenesis, or other means can be employed to obtain mutations of naturally occurring PKS-like genes to produce a polypeptide having PKS-like gene activity *in vivo* with more desirable

physical and kinetic parameters for function in the host cell, such as a longer half-life or a higher rate of production of a desired polyunsaturated fatty acid.

Of particular interest are the *Shewanella putrefaciens* ORFs and the corresponding ORFs of *Vibrio marinus*. The *Shewanella putrefaciens* PKS-like genes can be expressed in transgenic plants to effect biosynthesis of EPA. Other DNAs which are substantially identical in sequence to the *Shewanella putrefaciens* PKS-like genes, or which encode polypeptides which are substantially similar to PKS-like genes of *Shewanella putrefaciens* can be used, such as those identified from *Vibrio marinus*. By substantially identical in sequence is intended an amino acid sequence or nucleic acid sequence exhibiting in order of increasing preference at least 60%, 80%, 90% or 95% homology to the DNA sequence of the *Shewanella putrefaciens* PKS-like genes or nucleic acid sequences encoding the amino acid sequences for such genes. For polypeptides, the length of comparison sequences generally is at least 16 amino acids, preferably at least 20 amino acids, and most preferably 35 amino acids. For nucleic acids, the length of comparison sequences generally is at least 50 nucleotides, preferably at least 60 nucleotides, and more preferably at least 75 nucleotides, and most preferably, 110 nucleotides.

Homology typically is measured using sequence analysis software, for example, the Sequence Analysis software package of the Genetics Computer Group, University of Wisconsin Biotechnology Center, 1710 University Avenue, Madison, Wisconsin 53705, MEGAlign (DNAStar, Inc., 1228 S. Park St., Madison, Wisconsin 53715), and MacVector (Oxford Molecular Group, 2105 S. Bascom Avenue, Suite 200, Campbell, California 95008). BLAST (National Center for Biotechnology Information (NCBI) www.ncbi.nlm.nih.gov; FASTA (Pearson and Lipman, *Science* (1985) 227:1435-1446). Such software matches similar sequences by assigning degrees of homology to various substitutions, deletions, and other modifications. Conservative substitutions typically include substitutions within the following groups: glycine and alanine; valine, isoleucine and leucine; aspartic acid, glutamic acid, asparagine, and glutamine; serine and threonine; lysine and arginine; and phenylalanine and tyrosine. Substitutions may also be made on the basis of conserved hydrophobicity or hydrophilicity (Kyte and Doolittle, *J. Mol. Biol.* (1982) 157: 105-132), or on the basis of the ability to assume similar polypeptide secondary structure (Chou and Fasman, *Adv. Enzymol.* (1978) 47: 45-148, 1978). A

related protein to the probing sequence is identified when $p \geq 0.01$, preferably $p \geq 10^{-7}$ or 10^{-8} .

Encompassed by the present invention are related PKS-like genes from the same or other organisms. Such related PKS-like genes include variants of the disclosed PKS-like ORFs that occur naturally within the same or different species of *Shewanella*, as well as homologues of the disclosed PKS-like genes from other species and evolutionarily related proteins having analogous function and activity. Also included are PKS-like genes which, although not substantially identical to the *Shewanella putrefaciens* PKS-like genes, operate in a similar fashion to produce PUFAs as part of a PKS-like system.

Related PKS-like genes can be identified by their ability to function substantially the same as the disclosed PKS-like genes; that is, they can be substituted for corresponding ORFs of *Shewanella* or *Vibrio* and still effectively produce EPA or DHA. Related PKS-like genes also can be identified by screening sequence databases for sequences homologous to the disclosed PKS-like genes, by hybridization of a probe based on the disclosed PKS-like genes to a library constructed from the source organism, or by RT-PCR using mRNA from the source organism and primers based on the disclosed PKS-like gene. Thus, the phrase "PKS-like genes" refers not only to the nucleotide sequences disclosed herein, but also to other nucleic acids that are allelic or species variants of these nucleotide sequences. It is also understood that these terms include nonnatural mutations introduced by deliberate mutation using recombinant technology such as single site mutation or by excising short sections of DNA open reading frames coding for PUFA enzymes or by substituting new codons or adding new codons. Such minor alterations substantially maintain the immunoidentity of the original expression product and/or its biological activity. The biological properties of the altered PUFA enzymes can be determined by expressing the enzymes in an appropriate cell line and by determining the ability of the enzymes to synthesize PUFAs. Particular enzyme modifications considered minor would include substitution of amino acids of similar chemical properties, e.g., glutamic acid for aspartic acid or glutamine for asparagine.

When utilizing a PUFA PKS-like system from another organism, the regions of a PKS-like gene polypeptide important for PKS-like gene activity can be determined through routine mutagenesis, expression of the resulting mutant polypeptides and determination of their activities. The coding region for the mutants can include deletions, insertions and point mutations, or combinations thereof. A typical functional analysis

begins with deletion mutagenesis to determine the N- and C-terminal limits of the protein necessary for function, and then internal deletions, insertions or point mutants are made in the open ready frame to further determine regions necessary for function. Other techniques such as cassette mutagenesis or total synthesis also can be used. Deletion 5 mutagenesis is accomplished, for example, by using exonucleases to sequentially remove the 5' or 3' coding regions. Kits are available for such techniques. After deletion, the coding region is completed by ligating oligonucleotides containing start or stop codons to the deleted coding region after 5' or 3' deletion, respectively. Alternatively, oligonucleotides encoding start or stop codons are inserted into the coding region by a 10 variety of methods including site-directed mutagenesis, mutagenic PCR or by ligation onto DNA digested at existing restriction sites. Internal deletions can similarly be made through a variety of methods including the use of existing restriction sites in the DNA, by use of mutagenic primers via site directed mutagenesis or mutagenic PCR. Insertions are made through methods such as linker-scanning mutagenesis, site-directed mutagenesis or 15 mutagenic PCR. Point mutations are made through techniques such as site-directed mutagenesis or mutagenic PCR.

Chemical mutagenesis also can be used for identifying regions of a PKS-like gene polypeptide important for activity. A mutated construct is expressed, and the ability of the resulting altered protein to function as a PKS-like gene is assayed. Such structure- 20 function analysis can determine which regions may be deleted, which regions tolerate insertions, and which point mutations allow the mutant protein to function in substantially the same way as the native PKS-like gene. All such mutant proteins and nucleotide sequences encoding them are within the scope of the present invention. EPA is produced in *Shewanella* as the product of a PKS-like system, such that the EPA genes encode 25 components of this system. In *Vibrio*, DHA is produced by a similar system. The enzymes which synthesize these fatty acids are encoded by a cluster of genes which are distinct from the fatty acid synthesis genes encoding the enzymes involved in synthesis of the C16 and C18 fatty acids typically found in bacteria and in plants. As the *Shewanella* EPA genes represent a PKS-like gene cluster, EPA production is, at least to some extent, 30 independent of the typical bacterial type II FAS system. Thus, production of EPA in the cytoplasm of plant cells can be achieved by expression of the PKS-like pathway genes in plant cells under the control of appropriate plant regulatory signals.

EPA production in *E. coli* transformed with the *Shewanella* EPA genes proceeds during anaerobic growth, indicating that O₂-dependent desaturase reactions are not involved. Analyses of the proteins encoded by the ORFs essential for EPA production reveals the presence of domain structures characteristic of PKS-like systems. Fig. 2A 5 shows a summary of the domains, motifs, and also key homologies detected by "BLAST" data bank searches. Because EPA is different from many of the other substances produced by PKS-like pathways, i.e., it contains 5, *cis* double bonds, spaced at 3 carbon intervals along the molecule, a PKS-like system for synthesis of EPA is not expected.

Further, BLAST searches using the domains present in the *Shewanella* EPA ORFs 10 reveal that several are related to proteins encoded by a PKS-like gene cluster found in Anabeana. The structure of that region of the Anabeana chromosome is shown in Fig. 2F. The Anabeana PKS-like genes have been linked to the synthesis of a long-chain (C26), hydroxy-fatty acid found in a glycolipid layer of heterocysts. The EPA protein domains with homology to the Anabeana proteins are indicated in Fig. 2F.

15 ORF 6 of *Shewanella* contains a KAS domain which includes an active site motif (DXAC*) as well as a "GFGG" motif which is present at the end of many Type II KAS proteins (see Fig. 2A). Extended motifs are present but not shown here. Next is a malonyl-CoA:ACP acyl transferase (AT) domain. Sequences near the active site motif (GHS*XG) suggest it transfers malonate rather than methylmalonate, i.e., it resembles the 20 acetate-like ATs. Following a linker region, there is a cluster of 6 repeating domains, each ~100 amino acids in length, which are homologous to PKS-like ACP sequences. Each contains a pantetheine binding site motif (LGXDS*(L/I)). The presence of 6 such 25 ACP domains has not been observed previously in fatty acid synthases (FAS) or PKS-like systems. Near the end of the protein is a region which shows homology to β -keto-ACP reductases (KR). It contains a pyridine nucleotide binding site motif "GXGXX(G/A/P)".

The *Shewanella* ORF 8 begins with a KAS domain, including active site and 30 ending motifs (Fig. 2C). The best match in the data banks is with the Anabeana HglD. There is also a domain which has sequence homology to the N-terminal one half of the Anabeana HglC. This region also shows weak homology to KAS proteins although it lacks the active site and ending motifs. It has the characteristics of the so-called chain length factors (CLF) of Type II PKS-like systems. ORF 8 appears to direct the production of EPA versus DHA by the PKS-like system. ORF 8 also has two domains with homology to β -hydroxyacyl-ACP dehydrases (DH). The best match for both domains is

with *E. coli* FabA, a bi-functional enzyme which carries out both the dehydrase reaction and an isomerization (*trans* to *cis*) of the resulting double bond. The first DH domain contains both the active site histidine (H) and an adjacent cysteine (C) implicated in FabA catalysis. The second DH domain has the active site H but lacks the adjacent C (Fig. 2C).
5 Blast searches with the second DH domain also show matches to FabZ, a second *E. coli* DH, which does not possess isomerase activity.

The N-terminal half of ORF 7 (Fig. 2B) has no significant matches in the data banks. The best match of the C-terminal half is with a C-terminal portion of the Anabeana HglC. This domain contains an acyl-transferase (AT) motif (GX SXG).
10 Comparison of the extended active site sequences, based on the crystal structure of the *E. coli* malonyl-CoA:ACP AT, reveals that ORF 7 lacks two residues essential for exclusion of water from the active site (*E. coli* nomenclature; Q11 and R117). These data suggest that ORF 7 may function as a thioesterase.

ORF 9 (Fig. 2D) is homologous to an ORF of unknown function in the Anabeana Hgl cluster. It also exhibits a very weak homology to NIFA, a regulatory protein in nitrogen fixing bacteria. A regulatory role for the ORF 9 protein has not been excluded. ORF 3 (Fig. 2E) is homologous to the Anabeana HetI as well as EntD from *E. coli* and Sfp of *Bacillus*. Recently, a new enzyme family of phosphopantetheinyl transferases has been identified that includes HetI, EntD and Sfp (Lambot RH, *et al.* (1996) A new
20 enzyme superfamily - the phosphopantetheinyl transferases. *Chemistry & Biology*, Vol 3, #11, 923-936). The data of Fig. 3 demonstrates that the presence of ORF 3 is required for addition of β -alanine (i.e. pantetheine) to the ORF 6 protein. Thus, ORF 3 encodes the phosphopantetheinyl transferase specific for the ORF 6 ACP domains. (See, Haydock SF *et al.* (1995) Divergent sequence motifs correlated with the substrate specificity of
25 (methyl)malonyl-CoA:acyl carrier protein transacylase domains in modular polyketide synthases, *FEBS Lett.*, 374, 246-248). Malonate is the source of the carbons utilized in the extension reactions of EPA synthesis. Additionally, malonyl-CoA rather than malonyl-ACP is the AT substrate, i.e., the AT region of ORF 6 uses malonyl Co-A.

Once the DNA sequences encoding the PKS-like genes of an organism responsible
30 for PUFA production have been obtained, they are placed in a vector capable of replication in a host cell, or propagated *in vitro* by means of techniques such as PCR or long PCR. Replicating vectors can include plasmids, phage, viruses, cosmids and the like. Desirable vectors include those useful for mutagenesis of the gene of interest or for

expression of the gene of interest in host cells. A PUFA synthesis enzyme or a homologous protein can be expressed in a variety of recombinantly engineered cells. Numerous expression systems are available for expression of DNA encoding a PUFA enzyme. The expression of natural or synthetic nucleic acids encoding PUFA enzyme is typically achieved by operably linking the DNA to a promoter (which is either constitutive or inducible) within an expression vector. By expression vector is meant a DNA molecule, linear or circular, that comprises a segment encoding a PUFA enzyme, operably linked to additional segments that provide for its transcription. Such additional segments include promoter and terminator sequences. An expression vector also may include one or more origins of replication, one or more selectable markers, an enhancer, a polyadenylation signal, etc. Expression vectors generally are derived from plasmid or viral DNA, and can contain elements of both. The term "operably linked" indicates that the segments are arranged so that they function in concert for their intended purposes, for example, transcription initiates in the promoter and proceeds through the coding segment to the terminator. *See Sambrook et al, supra.*

The technique of long PCR has made *in vitro* propagation of large constructs possible, so that modifications to the gene of interest, such as mutagenesis or addition of expression signals, and propagation of the resulting constructs can occur entirely *in vitro* without the use of a replicating vector or a host cell. *In vitro* expression can be accomplished, for example, by placing the coding region for the desaturase polypeptide in an expression vector designed for *in vitro* use and adding rabbit reticulocyte lysate and cofactors; labeled amino acids can be incorporated if desired. Such *in vitro* expression vectors may provide some or all of the expression signals necessary in the system used. These methods are well known in the art and the components of the system are commercially available. The reaction mixture can then be assayed directly for PKS-like enzymes for example by determining their activity, or the synthesized enzyme can be purified and then assayed.

Expression in a host cell can be accomplished in a transient or stable fashion. Transient expression can occur from introduced constructs which contain expression signals functional in the host cell, but which constructs do not replicate and rarely integrate in the host cell, or where the host cell is not proliferating. Transient expression also can be accomplished by inducing the activity of a regulatable promoter operably linked to the gene of interest, although such inducible systems frequently exhibit a low

basal level of expression. Stable expression can be achieved by introduction of a nucleic acid construct that can integrate into the host genome or that autonomously replicates in the host cell. Stable expression of the gene of interest can be selected for through the use of a selectable marker located on or transfected with the expression construct, followed by 5 selection for cells expressing the marker. When stable expression results from integration, integration of constructs can occur randomly within the host genome or can be targeted through the use of constructs containing regions of homology with the host genome sufficient to target recombination with the host locus. Where constructs are targeted to an endogenous locus, all or some of the transcriptional and translational 10 regulatory regions can be provided by the endogenous locus. To achieve expression in a host cell, the transformed DNA is operably associated with transcriptional and translational initiation and termination regulatory regions that are functional in the host cell.

Transcriptional and translational initiation and termination regions are derived 15 from a variety of nonexclusive sources, including the DNA to be expressed, genes known or suspected to be capable of expression in the desired system, expression vectors, chemical synthesis. The termination region can be derived from the 3' region of the gene from which the initiation region was obtained or from a different gene. A large number 20 of termination regions are known to and have been found to be satisfactory in a variety of hosts from the same and different genera and species. The termination region usually is selected more as a matter of convenience rather than because of any particular property. When expressing more than one PKS-like ORF in the same cell, appropriate regulatory 25 regions and expression methods should be used. Introduced genes can be propagated in the host cell through use of replicating vectors or by integration into the host genome. Where two or more genes are expressed from separate replicating vectors, it is desirable 30 that each vector has a different means of replication. Each introduced construct, whether integrated or not, should have a different means of selection and should lack homology to the other constructs to maintain stable expression and prevent reassortment of elements among constructs. Judicious choices of regulatory regions, selection means and method of propagation of the introduced construct can be experimentally determined so that all introduced genes are expressed at the necessary levels to provide for synthesis of the desired products.

A variety of prokaryotic expression systems can be used to express PUFA enzyme. Expression vectors can be constructed which contain a promoter to direct transcription, a ribosome binding site, and a transcriptional terminator. Examples of regulatory regions suitable for this purpose in *E. coli* are the promoter and operator region of the *E. coli* tryptophan biosynthetic pathway as described by Yanofsky (1984) *J. Bacteriol.*, 158:1018-1024 and the leftward promoter of phage lambda (P λ) as described by Herskowitz and Hagen, (1980) *Ann. Rev. Genet.*, 14:399-445. The inclusion of selection markers in DNA vectors transformed in *E. coli* is also useful. Examples of such markers include genes specifying resistance to ampicillin, tetracycline, or chloramphenicol.

Vectors used for expressing foreign genes in bacterial hosts generally will contain a selectable marker, such as a gene for antibiotic resistance, and a promoter which functions in the host cell. Plasmids useful for transforming bacteria include pBR322 (Bolivar, *et al*, (1977) *Gene* 2:95-113), the pUC plasmids (Messing, (1983) *Meth. Enzymol.* 101:20-77, Vieira and Messing, (1982) *Gene* 19:259-268), pCQV2 (Queen, *ibid.*), and derivatives thereof. Plasmids may contain both viral and bacterial elements. Methods for the recovery of the proteins in biologically active form are discussed in U.S. Patent Nos. 4,966,963 and 4,999,422, which are incorporated herein by reference. See Sambrook, *et al* for a description of other prokaryotic expression systems.

For expression in eukaryotes, host cells for use in practicing the present invention include mammalian, avian, plant, insect, and fungal cells. As an example, for plants, the choice of a promoter will depend in part upon whether constitutive or inducible expression is desired and whether it is desirable to produce the PUFAAs at a particular stage of plant development and/or in a particular tissue. Considerations for choosing a specific tissue and/or developmental stage for expression of the ORFs may depend on competing substrates or the ability of the host cell to tolerate expression of a particular PUFA. Expression can be targeted to a particular location within a host plant such as seed, leaves, fruits, flowers, and roots, by using specific regulatory sequences, such as those described in USPN 5,463,174, USPN 4,943,674, USPN 5,106,739, USPN 5,175,095, USPN 5,420,034, USPN 5,188,958, and USPN 5,589,379. Where the host cell is a yeast, transcription and translational regions functional in yeast cells are provided, particularly from the host species. The transcriptional initiation regulatory regions can be obtained, for example from genes in the glycolytic pathway, such as alcohol dehydrogenase, glyceraldehyde-3-phosphate dehydrogenase (GPD),

phosphoglucoisomerase, phosphoglycerate kinase, etc. or regulatable genes such as acid phosphatase, lactase, metallothionein, glucoamylase, etc. Any one of a number of regulatory sequences can be used in a particular situation, depending upon whether constitutive or induced transcription is desired, the particular efficiency of the promoter in conjunction with the open-reading frame of interest, the ability to join a strong promoter with a control region from a different promoter which allows for inducible transcription, ease of construction, and the like. Of particular interest are promoters which are activated in the presence of galactose. Galactose-inducible promoters (GAL1, GAL7, and GAL10) have been extensively utilized for high level and regulated expression of protein in yeast (Lue *et al.*, (1987) *Mol. Cell. Biol.* 7:3446; Johnston, (1987) *Microbiol. Rev.* 51:458). Transcription from the GAL promoters is activated by the GAL4 protein, which binds to the promoter region and activates transcription when galactose is present. In the absence of galactose, the antagonist GAL80 binds to GAL4 and prevents GAL4 from activating transcription. Addition of galactose prevents GAL80 from inhibiting activation by GAL4. Preferably, the termination region is derived from a yeast gene, particularly *Saccharomyces*, *Schizosaccharomyces*, *Candida* or *Kluyveromyces*. The 3' regions of two mammalian genes, γ interferon and α 2 interferon, are also known to function in yeast.

Nucleotide sequences surrounding the translational initiation codon ATG have been found to affect expression in yeast cells. If the desired polypeptide is poorly expressed in yeast, the nucleotide sequences of exogenous genes can be modified to include an efficient yeast translation initiation sequence to obtain optimal gene expression. For expression in *Saccharomyces*, this can be done by site-directed mutagenesis of an inefficiently expressed gene by fusing it in-frame to an endogenous *Saccharomyces* gene, preferably a highly expressed gene, such as the lactase gene.

As an alternative to expressing the PKS-like genes in the plant cell cytoplasm, is to target the enzymes to the chloroplast. One method to target proteins to the chloroplast entails use of leader peptides attached to the N-termini of the proteins. Commonly used leader peptides are derived from the small subunit of plant ribulose bis phosphate carboxylase. Leader sequences from other chloroplast proteins may also be used. Another method for targeting proteins to the chloroplast is to transform the chloroplast genome (Stable transformation of chloroplasts of *Chlamydomonas reinhardtii* (1 green alga) using bombardment of recipient cells with high-velocity tungsten microprojectiles coated with foreign DNA has been described. *See*, for example, Blowers *et al* *Plant Cell*

(1989) 1:123-132 and Debuchy *et al* *EMBO J* (1989) 8:2803-2809. The transformation technique, using tungsten microprojectiles, is described by Kline *et al*, *Nature* (London) (1987) 327:70-73). The most common method of transforming chloroplasts involves using biolistic techniques, but other techniques developed for the purpose may also be used. (Methods for targeting foreign gene products into chloroplasts (Shrier *et al* *EMBO J.* (1985) 4:25-32) or mitochondria (Boutry *et al*, *supra*) have been described. See also Tomai *et al* *Gen. Biol. Chem.* (1988) 263:15104-15109 and US Patent No. 4,940,835 for the use of transit peptides for translocating nuclear gene products into the chloroplast. Methods for directing the transport of proteins to the chloroplast are reviewed in Kenauf *TIBTECH* (1987) 5:40-47.

For producing PUFAs in avian species and cells, gene transfer can be performed by introducing a nucleic acid sequence encoding a PUFA enzyme into the cells following procedures known in the art. If a transgenic animal is desired, pluripotent stem cells of embryos can be provided with a vector carrying a PUFA enzyme encoding transgene and developed into adult animal (USPN 5,162,215; Ono *et al.* (1996) *Comparative Biochemistry and Physiology A* 113(3):287-292; WO 9612793; WO 9606160). In most cases, the transgene is modified to express high levels of the PKS-like enzymes in order to increase production of PUFAs. The transgenes can be modified, for example, by providing transcriptional and/or translational regulatory regions that function in avian cells, such as promoters which direct expression in particular tissues and egg parts such as yolk. The gene regulatory regions can be obtained from a variety of sources, including chicken anemia or avian leukosis viruses or avian genes such as a chicken ovalbumin gene.

Production of PUFAs in insect cells can be conducted using baculovirus expression vectors harboring PKS-like transgenes. Baculovirus expression vectors are available from several commercial sources such as Clonetech. Methods for producing hybrid and transgenic strains of algae, such as marine algae, which contain and express a desaturase transgene also are provided. For example, transgenic marine algae can be prepared as described in USPN 5,426,040. As with the other expression systems described above, the timing, extent of expression and activity of the desaturase transgene can be regulated by fitting the polypeptide coding sequence with the appropriate transcriptional and translational regulatory regions selected for a particular use. Of particular interest are promoter regions which can be induced under preselected growth

conditions. For example, introduction of temperature sensitive and/or metabolite responsive mutations into the desaturase transgene coding sequences, its regulatory regions, and/or the genome of cells into which the transgene is introduced can be used for this purpose.

5 The transformed host cell is grown under appropriate conditions adapted for a desired end result. For host cells grown in culture, the conditions are typically optimized to produce the greatest or most economical yield of PUFAs, which relates to the selected desaturase activity. Media conditions which may be optimized include: carbon source, nitrogen source, addition of substrate, final concentration of added substrate, form of 10 substrate added, aerobic or anaerobic growth, growth temperature, inducing agent, induction temperature, growth phase at induction, growth phase at harvest, pH, density, and maintenance of selection. Microorganisms such as yeast, for example, are preferably grown using selected media of interest, which include yeast peptone broth (YPD) and minimal media (contains amino acids, yeast nitrogen base, and ammonium sulfate, and 15 lacks a component for selection, for example uracil). Desirably, substrates to be added are first dissolved in ethanol. Where necessary, expression of the polypeptide of interest may be induced, for example by including or adding galactose to induce expression from a GAL promoter.

When increased expression of the PKS-like gene polypeptide in a host cell which 20 expresses PUFA from a PKS-like system is desired, several methods can be employed. Additional genes encoding the PKS-like gene polypeptide can be introduced into the host organism. Expression from the native PKS-like gene locus also can be increased through homologous recombination, for example by inserting a stronger promoter into the host genome to cause increased expression, by removing destabilizing sequences from either 25 the mRNA or the encoded protein by deleting that information from the host genome, or by adding stabilizing sequences to the mRNA (*see USPN 4,910,141 and USPN 5,500,365*). Thus, the subject host will have at least have one copy of the expression construct and may have two or more, depending upon whether the gene is integrated into the genome, amplified, or is present on an extrachromosomal element having multiple 30 copy numbers. Where the subject host is a yeast, four principal types of yeast plasmid vectors can be used: Yeast Integrating plasmids (YIps), Yeast Replicating plasmids (YRps), Yeast Centromere plasmids (YCps), and Yeast Episomal plasmids (YEps). YIps lack a yeast replication origin and must be propagated as integrated elements in the yeast

genome. YRps have a chromosomally derived autonomously replicating sequence and are propagated as medium copy number (20 to 40), autonomously replicating, unstably segregating plasmids. YCps have both a replication origin and a centromere sequence and propagate as low copy number (10-20), autonomously replicating, stably segregating plasmids. YEps have an origin of replication from the yeast 2 μ m plasmid and are propagated as high copy number, autonomously replicating, irregularly segregating plasmids. The presence of the plasmids in yeast can be ensured by maintaining selection for a marker on the plasmid. Of particular interest are the yeast vectors pYES2 (a YEplasmid available from Invitrogen, confers uracil prototrophy and a GAL1 galactose-inducible promoter for expression), and pYX424 (a YEplasmid having a constitutive TP1 promoter and conferring leucine prototrophy; (Alber and Kawasaki (1982). *J. Mol. & Appl. Genetics* 1: 419).

The choice of a host cell is influenced in part by the desired PUFA profile of the transgenic cell, and the native profile of the host cell. Even where the host cell expresses PKS-like gene activity for one PUFA, expression of PKS-like genes of another PKS-like system can provide for production of a novel PUFA not produced by the host cell. In particular instances where expression of PKS-like gene activity is coupled with expression of an ORF 8 PKS-like gene of an organism which produces a different PUFA, it can be desirable that the host cell naturally have, or be mutated to have, low PKS-like gene activity for ORF 8. As an example, for production of EPA, the DNA sequence used encodes the polypeptide having PKS-like gene activity of an organism which produces EPA, while for production of DHA, the DNA sequences used are those from an organism which produces DHA. For use in a host cell which already expresses PKS-like gene activity it can be necessary to utilize an expression cassette which provides for overexpression of the desired PKS-like genes alone or with a construct to downregulate the activity of an existing ORF of the existing PKS-like system, such as by antisense or co-suppression. Similarly, a combination of ORFs derived from separate organisms which produce the same or different PUFAAs using PKS-like systems may be used. For instance, the ORF 8 of *Vibrio* directs the expression of DHA in a host cell, even when ORFs 3, 6, 7 and 9 are from *Shewanella*, which produce EPA when coupled to ORF 8 of *Shewanella*. Therefore, for production of eicosapentanoic acid (EPA), the expression cassettes used generally include one or more cassettes which include ORFs 3, 6, 7, 8 and 9 from a PUFA-producing organism such as the marine bacterium *Shewanella*.

putrefaciens (for EPA production) or *Vibrio marinus* (for DHA production). ORF 8 can be used for induction of DHA production, and ORF 8 of *Vibrio* can be used in conjunction with ORFs 3, 6, 7 and 9 of *Shewanella* to produce DHA. The organization and numbering scheme of the ORFs identified in the *Shewanella* gene cluster are shown 5 in Fig 1A. Maps of several subclones referred to in this study are shown in Fig 1B. For expression of a PKS-like gene polypeptide, transcriptional and translational initiation and termination regions functional in the host cell are operably linked to the DNA encoding the PKS-like gene polypeptide.

Constructs comprising the PKS-like ORFs of interest can be introduced into a host 10 cell by any of a variety of standard techniques, depending in part upon the type of host cell. These techniques include transfection, infection, bolistic impact, electroporation, microinjection, scraping, or any other method which introduces the gene of interest into the host cell (see USPN 4,743,548, USPN 4,795,855, USPN 5,068,193, USPN 5,188,958, USPN 5,463,174, USPN 5,565,346 and USPN 5,565,347). Methods of transformation 15 which are used include lithium acetate transformation (*Methods in Enzymology*, (1991) 194:186-187). For convenience, a host cell which has been manipulated by any method to take up a DNA sequence or construct will be referred to as "transformed" or "recombinant" herein. The subject host will have at least have one copy of the expression construct and may have two or more, depending upon whether the gene is integrated into 20 the genome, amplified, or is present on an extrachromosomal element having multiple copy numbers.

For production of PUFAs, depending upon the host cell, the several polypeptides produced by pEPA, ORFs 3, 6, 7, 8 and 9, are introduced as individual expression constructs or can be combined into two or more cassettes which are introduced 25 individually or co-transformed into a host cell. A standard transformation protocol is used. For plants, where less than all PKS-like genes required for PUFA synthesis have been inserted into a single plant, plants containing a complementing gene or genes can be crossed to obtain plants containing a full complement of PKS-like genes to synthesize a desired PUFA.

30 The PKS-like-mediated production of PUFAs can be performed in either prokaryotic or eukaryotic host cells. The cells can be cultured or formed as part or all of a host organism including an animal. Viruses and bacteriophage also can be used with appropriate cells in the production of PUFAs, particularly for gene transfer, cellular

targeting and selection. Any type of plant cell can be used for host cells, including dicotyledonous plants, monocotyledonous plants, and cereals. Of particular interest are crop plants such as *Brassica*, *Arabidopsis*, soybean, corn, and the like. Prokaryotic cells of interest include *Escherichia*, *Bacillus*, *Lactobacillus*, *cyanobacteria* and the like.

5 Eukaryotic cells include plant cells, mammalian cells such as those of lactating animals, avian cells such as of chickens, and other cells amenable to genetic manipulation including insect, fungal, and algae cells. Examples of host animals include mice, rats, rabbits, chickens, quail, turkeys, cattle, sheep, pigs, goats, yaks, etc., which are amenable to genetic manipulation and cloning for rapid expansion of a transgene expressing
10 population. For animals, PKS-like transgenes can be adapted for expression in target organelles, tissues and body fluids through modification of the gene regulatory regions. Of particular interest is the production of PUFAs in the breast milk of the host animal.

Examples of host microorganisms include *Saccharomyces cerevisiae*, *Saccharomyces carlsbergensis*, or other yeast such as *Candida*, *Kluyveromyces* or other fungi, for example, filamentous fungi such as *Aspergillus*, *Neurospora*, *Penicillium*, etc. Desirable characteristics of a host microorganism are, for example, that it is genetically well characterized, can be used for high level expression of the product using ultra-high density fermentation, and is on the GRAS (generally recognized as safe) list since the proposed end product is intended for ingestion by humans. Of particular interest is use of
20 a yeast, more particularly baker's yeast (*S. cerevisiae*), as a cell host in the subject invention. Strains of particular interest are SC334 (Mat α pep4-3 prbl-1122 ura3-52 leu2-3, 112 regl-501 gall1; (Hovland *et al* (1989) Gene 83:57-64); BJ1995 (Yeast Genetic Stock Centre, 1021 Donner Laboratory, Berkeley, CA 94720), INVSC1 (Mat α hiw3 Δ 1 leu2 trp1-289 ura3-52 (Invitrogen, 1600 Faraday Ave., Carlsbad, CA 92008) and INVSC2
25 (Mat α his3 Δ 200 ura3-167; (Invitrogen). Bacterial cells also may be used as hosts. This includes *E. coli*, which can be useful in fermentation processes. Alternatively, a host such as a *Lactobacillus* species can be used as a host for introducing the products of the PKS-like pathway into a product such as yogurt.

30 The transformed host cell can be identified by selection for a marker contained on the introduced construct. Alternatively, a separate marker construct can be introduced with the desired construct, as many transformation techniques introduce multiple DNA molecules into host cells. Typically, transformed hosts are selected for their ability to grow on selective media. Selective media can incorporate an antibiotic or lack a factor

necessary for growth of the untransformed host, such as a nutrient or growth factor. An introduced marker gene therefor may confer antibiotic resistance, or encode an essential growth factor or enzyme, and permit growth on selective media when expressed in the transformed host cell. Desirably, resistance to kanamycin and the amino glycoside G418 are of particular interest (see USPN 5,034,322). For yeast transformants, any marker that functions in yeast can be used, such as the ability to grow on media lacking uracil, leucine, lysine or tryptophan.

Selection of a transformed host also can occur when the expressed marker protein can be detected, either directly or indirectly. The marker protein can be expressed alone or as a fusion to another protein. The marker protein can be one which is detected by its enzymatic activity; for example β -galactosidase can convert the substrate X-gal to a colored product, and luciferase can convert luciferin to a light-emitting product. The marker protein can be one which is detected by its light-producing or modifying characteristics; for example, the green fluorescent protein of *Aequorea victoria* fluoresces when illuminated with blue light. Antibodies can be used to detect the marker protein or a molecular tag on, for example, a protein of interest. Cells expressing the marker protein or tag can be selected, for example, visually, or by techniques such as FACS or panning using antibodies.

The PUFAs produced using the subject methods and compositions are found in the host plant tissue and/or plant part as free fatty acids and/or in conjugated forms such as acylglycerols, phospholipids, sulfolipids or glycolipids, and can be extracted from the host cell through a variety of means well-known in the art. Such means include extraction with organic solvents, sonication, supercritical fluid extraction using for example carbon dioxide, and physical means such as presses, or combinations thereof. Of particular interest is extraction with methanol and chloroform. Where appropriate, the aqueous layer can be acidified to protonate negatively charged moieties and thereby increase partitioning of desired products into the organic layer. After extraction, the organic solvents can be removed by evaporation under a stream of nitrogen. When isolated in conjugated forms, the products are enzymatically or chemically cleaved to release the free fatty acid or a less complex conjugate of interest, and are then subjected to further manipulations to produce a desired end product. Desirably, conjugated forms of fatty acids are cleaved with potassium hydroxide.

If further purification is necessary, standard methods can be employed. Such methods include extraction, treatment with urea, fractional crystallization, HPLC, fractional distillation, silica gel chromatography, high speed centrifugation or distillation, or combinations of these techniques. Protection of reactive groups, such as the acid or 5 alkenyl groups, can be done at any step through known techniques, for example alkylation or iodination. Methods used include methylation of the fatty acids to produce methyl esters. Similarly, protecting groups can be removed at any step. Desirably, purification of fractions containing DHA and EPA is accomplished by treatment with urea and/or fractional distillation.

10 The uses of the subject invention are several. Probes based on the DNAs of the present invention find use in methods for isolating related molecules or in methods to detect organisms expressing PKS-like genes. When used as probes, the DNAs or oligonucleotides need to be detectable. This is usually accomplished by attaching a label either at an internal site, for example via incorporation of a modified residue, or at the 5' 15 or 3' terminus. Such labels can be directly detectable, can bind to a secondary molecule that is detectably labeled, or can bind to an unlabelled secondary molecule and a detectably labeled tertiary molecule; this process can be extended as long as is practicable to achieve a satisfactorily detectable signal without unacceptable levels of background signal. Secondary, tertiary, or bridging systems can include use of antibodies directed 20 against any other molecule, including labels or other antibodies, or can involve any molecules which bind to each other, for example a biotin-streptavidin/avidin system. Detectable labels typically include radioactive isotopes, molecules which chemically or enzymatically produce or alter light, enzymes which produce detectable reaction products, magnetic molecules, fluorescent molecules or molecules whose fluorescence or light- 25 emitting characteristics change upon binding. Examples of labelling methods can be found in USPN 5,011,770. Alternatively, the binding of target molecules can be directly detected by measuring the change in heat of solution on binding of a probe to a target via isothermal titration calorimetry, or by coating the probe or target on a surface and detecting the change in scattering of light from the surface produced by binding of a target 30 or a probe, respectively, is done with the BIACore system.

PUFAs produced by recombinant means find applications in a wide variety of areas. Supplementation of humans or animals with PUFAs in various forms can result in increased levels not only of the added PUFAs, but of their metabolic progeny as well.

Complex regulatory mechanisms can make it desirable to combine various PUFAs, or to add different conjugates of PUFAs, in order to prevent, control or overcome such mechanisms to achieve the desired levels of specific PUFAs in an individual. In the present case, expression of PKS-like gene genes, or antisense PKS-like gene transcripts, 5 can alter the levels of specific PUFAs, or derivatives thereof, found in plant parts and/or plant tissues. The PKS-like gene polypeptide coding region is expressed either by itself or with other genes, in order to produce tissues and/or plant parts containing higher proportions of desired PUFAs or containing a PUFA composition which more closely resembles that of human breast milk (Prieto *et al.*, PCT publication WO 95/24494) than 10 does the unmodified tissues and/or plant parts.

PUFAs, or derivatives thereof, made by the disclosed method can be used as dietary supplements for patients undergoing intravenous feeding or for preventing or treating malnutrition. For dietary supplementation, the purified PUFAs, or derivatives thereof, can be incorporated into cooking oils, fats or margarines formulated so that in 15 normal use the recipient receives a desired amount of PUFA. The PUFAs also can be incorporated into infant formulas, nutritional supplements or other food products, and find use as anti-inflammatory or cholesterol lowering agents.

Particular fatty acids such as EPA can be used to alter the composition of infant formulas to better replicate the PUFA composition of human breast milk. The 20 predominant triglyceride in human milk is reported to be 1,3-di-oleoyl-2-palmitoyl, with 2-palmitoyl glycerides reported as better absorbed than 2-oleoyl or 2-linoleyl glycerides (*see* USPN 4,876,107). Typically, human breast milk has a fatty acid profile comprising from about 0.15 % to about 0.36 % as DHA, from about 0.03 % to about 0.13 % as EPA, from about 0.30 % to about 0.88 % as ARA, from about 0.22 % to about 0.67 % as 25 DGLA, and from about 0.27 % to about 1.04 % as GLA. A preferred ratio of GLA:DGLA:ARA in infant formulas is from about 1:1:4 to about 1:1:1, respectively. Amounts of oils providing these ratios of PUFA can be determined without undue experimentation by one of skill in the art. PUFAs, or host cells containing them, also can 30 be used as animal food supplements to alter an animal's tissue or milk fatty acid composition to one more desirable for human or animal consumption.

For pharmaceutical use (human or veterinary), the compositions generally are administered orally but can be administered by any route by which they may be successfully absorbed, e.g., parenterally (i.e. subcutaneously, intramuscularly or

intravenously), rectally or vaginally or topically, for example, as a skin ointment or lotion. Where available, gelatin capsules are the preferred form of oral administration. Dietary supplementation as set forth above also can provide an oral route of administration. The unsaturated acids of the present invention can be administered in conjugated forms, or as 5 salts, esters, amides or prodrugs of the fatty acids. Any pharmaceutically acceptable salt is encompassed by the present invention; especially preferred are the sodium, potassium or lithium salts. Also encompassed are the N-alkylpolyhydroxamine salts, such as N-methyl glucamine, described in PCT publication WO 96/33155. Preferred esters are the ethyl esters.

10 The PUFAs of the present invention can be administered alone or in combination with a pharmaceutically acceptable carrier or excipient. As solid salts, the PUFAs can also be administered in tablet form. For intravenous administration, the PUFAs or derivatives thereof can be incorporated into commercial formulations such as Intralipids. Where desired, the individual components of formulations can be individually provided in 15 kit form, for single or multiple use. A typical dosage of a particular fatty acid is from 0.1 mg to 20 g, or even 100 g daily, and is preferably from 10 mg to 1, 2, 5 or 10 g daily as required, or molar equivalent amounts of derivative forms thereof. Parenteral nutrition compositions comprising from about 2 to about 30 weight percent fatty acids calculated as triglycerides are encompassed by the present invention. Other vitamins, and 20 particularly fat-soluble vitamins such as vitamin A, D, E and L-carnitine optionally can be included. Where desired, a preservative such as a tocopherol can be added, typically at about 0.1% by weight.

The following examples are presented by way of illustration, not of limitation.

25

EXAMPLES

Example 1

The Identity of ORFs Derived from *Vibrio marinus*

Using polymerase chain reaction (PCR) with primers based on ORF 6 of 30 *Shewanella* (Sp ORF 6) sequences (FW 5' primers CUACUACUACUACCAAGCT AAAGCACTTAACCGTG, and CUACUACUACUAACAGCGAAATGCTTATCAAG for *Vibrio* and SS9 respectively and 3' BW primers: CAUCAUCAUCAUGCGACC

AAAACCAAATGAGCTAATAC for both *Vibrio* and SS9) and genomic DNAs templates from *Vibrio* and a borophyllic *photobacter* producing EPA (provided by Dr. Bartlett, UC San Diego), resulted in PCR products of *ca.*400 bases for *Vibrio marinus* (*Vibrio*) and *ca.*900 bases for SS9 presenting more than 75% homology with 5 corresponding fragments of Sp ORF 6 (see Figure 25) as determined by direct counting of homologous amino acids.

A *Vibrio* cosmid library was then prepared and using the *Vibrio* ORF 6 PCR product as a probe (see Figure 26); clones containing at least ORF 6 were selected by colony hybridization.

10 Through additional sequences of the selected cosmids such as cosmid #9 and cosmid #21, a *Vibrio* cluster (Figure 5) with ORFs homologous to, and organized in the same sequential order (ORFs 6-9) as ORFs 6-9 of *Shewanella*, was obtained (Figure 7). The *Vibrio* ORFs from this sequence are found at 17394 to 36115 and comprehend ORFs 6-9.

15

Table
Vibrio operon figures

	17394 to 25349	length = 7956 nt
	25509 to 28157	length = 2649 nt
20	28209 to 34262	length = 6054 nt
	34454 to 36115	length = 1662 nt

The ORF designations for the *Shewanella* genes are based on those disclosed in Figure 4, and differ from those published for the *Shewanella* cluster (Yazawa *et al*, USPN 25 5,683,898). For instance, ORF 3 of Figure 4 is read in the opposite direction from the other ORFs and is not disclosed in Yazawa *et al* USPN 5,683,898 (See Fig. 24) for comparison with Yazawa *et al* USPN 5,683,898).

Sequences homologous to ORF 3, were not found in the proximity of ORF 6 (17000 bases upstream of ORF 6) or of ORF 9 (*ca.*4000 bases downstream of ORF 9). 30 Motifs characteristic of phosphopantethenyl transferases (Lambalot *et al* (1996) *Current Biology* 3:923-936) were absent from the *Vibrio* sequences screened for these motifs. In addition, there was no match to Sp ORF 3 derived probes in genomic digests of *Vibrio* and of SC2A *Shewanella* (another bacterium provided by the University of San Diego and

also capable of producing EPA). Although ORF 3 may exist in *Vibrio*, its DNA may not be homologous to that of *Sp* ORF 3 and/or could be located in portions of the genome that were not sequenced.

Figure 6 provides the sequence of an approximately 19 kb *Vibrio* clone comprising ORFs 6-9. Figures 7 and 8 compare the gene cluster organizations of the PKS-like systems of *Vibrio marinus* and *Shewanella putrefaciens*. Figures 9 through 12 show the levels of sequence homology between the corresponding ORFs 6, 7, 8 and 9, respectively.

Example 2

ORF 8 Directs DHA Production

As described in example 1, DNA homologous to *Sp* ORF 6 was found in an unrelated species, SS9 *Photobacter*, which also is capable of producing EPA. Additionally, ORFs homologous to *Sp* ORF 6-9 were found in the DHA producing *Vibrio marinus* (*Vibrio*). From these ORFs a series of experiments was designed in which deletions in each of *Sp* ORFs 6-9 that suppressed EPA synthesis in *E. coli* (Yazawa (1996) *supra*) were complemented by the corresponding homologous genes from *Vibrio*.

The *Sp* EPA cluster was used to determine if any of the *Vibrio* ORFs 6-9 was responsible for the production of DHA. Deletion mutants provided for each of the *Sp* ORFs are EPA and DHA null. Each deletion was then complemented by the corresponding *Vibrio* ORF expressed behind a *lac* promoter (Figure 13).

The complementation of a *Sp* ORF 6 deletion by a *Vibrio* ORF 6 reestablished the production of EPA. Similar results were obtained by complementing the *Sp* ORF 7 and ORF 9 deletions. By contrast, the complementation of a *Sp* ORF 8 deletion resulted in the production of C22:6. *Vibrio* ORF 8 therefore appears to be a key element in the synthesis of DHA. Figures 14 and 15 show chromatograms of fatty acid profiles from the respective complementations of *Sp* del ORF 6 with *Vibrio* ORF 6 (EPA and no DHA) and *Sp* del ORF 8 with *Vibrio* ORF 8 (DHA). Figure 16 shows the fatty acid percentages for the ORF 8 complementation, again demonstrating that ORF 8 is responsible for DHA production.

These data show that polyketide-like synthesis genes with related or similar ORFs can be combined and expressed in a heterologous system and used to produce a distinct PUFA species in the host system, and that ORF 8 has a role in determining the ultimate chain length. The *Vibrio* ORFs 6, 7, 8, and 9 reestablish EPA synthesis. In the case of

Vibrio ORF 8, DHA is also present (*ca.* 0.7%) along with EPA (*ca.* 0.6%) indicating that this gene plays a significant role in directing synthesis of DHA *vs* EPA for these systems.

Example 3

5

Requirements for Production of DHA

To determine how *Vibrio* ORFs of the cluster ORF 6-9 are used in combination with *Vibrio* ORF 8, some combinations of *Vibrio* ORF 8 with some or all of the other *Vibrio* ORFs 6-9 cluster were created to explain the synthesis of DHA.

10 *Vibrio* ORFs 6-9 were complemented with *Sp* ORF 3. The results of this complementation are presented in Figures 16b and 16c. The significant amounts of DHA measured (greater than about 9%) and the absence of EPA suggest that no ORFs other than those of *Vibrio* ORFs 6-9 are required for DHA synthesis when combined with *Sp* ORF 3. This suggests that *Sp* ORF 3 plays a general function in the synthesis of bacterial PUFAs.

15 With respect to the DHA *vs* EPA production, it may be necessary to combine *Vibrio* ORF 8 with other *Vibrio* ORFs of the 6-9 cluster in order to specifically produce DHA. The roles of *Vibrio* ORF 9 and each of the combinations of *Vibrio* ORFs (6,8), (7, 8), (8, 9), etc in the synthesis of DHA are being studied.

20

Example 4

Plant Expression Constructs

A cloning vector with very few restriction sites was designed to facilitate the cloning of large fragments and their subsequent manipulation. An adapter was assembled by annealing oligonucleotides with the sequences AAGCCCGGGCTT and 25 GTACAAGCCCGGGCTTAGCT. This adapter was ligated to the vector pBluescript II SK+ (Stratagene) after digestion of the vector with the restriction endonucleases *Asp*718 and *Sst*I. The resulting vector, pCGN7769 had a single *Srf*I (and embedded *Sma*I) cloning site for the cloning of blunt ended DNA fragments.

30 A plasmid containing the napin cassette from pCGN3223, (USPN 5,639,790) was modified to make it more useful for cloning large DNA fragments containing multiple restriction sites, and to allow the cloning of multiple napin fusion genes into plant binary transformation vectors. An adapter comprised of the self annealed oligonucleotide of sequence CGCGATTAAATGGCGCGCCCTGCAGGCAGGCCCTGCAGGGCGC

GCCATTAAAT was ligated into the vector pBC SK+ (Stratagene) after digestion of the vector with the restriction endonuclease *Bss*HII to construct vector pCGN7765. Plamids pCGN3223 and pCGN7765 were digested with *Not*I and ligated together. The resultant vector, pCGN7770 (Figure 17), contains the pCGN7765 backbone and the napin seed specific expression cassette from pCGN3223.

Shewanella constructs

Genes encoding the *Shewanella* proteins were mutagenized to introduce suitable cloning sites 5' and 3' ORFs using PCR. The template for the PCR reactions was DNA of the cosmid pEPA (Yazawa *et al, supra*). PCR reactions were performed using Pfu DNA polymerase according to the manufacturers' protocols. The PCR products were cloned into *Srf*I digested pCGN7769. The primers CTGCAGCTCGAGACAATGTTGATT TCCTTATACCTCTGTCC and GGATCCAGATCTCTAGCTAGCTAGCTGAAGC TCGA were used to amplify ORF 3, and to generate plasmid pCGN8520. The primers 10 TCTAGACTCGAGACAATGAGGCCAGACCTCTAAACCTACA and CCCGGGCTC GAGCTAATTGCCCTCACTGTCGTTGCT were used to amplify ORF 6, and generate plasmid pCGN7776. The primers GAATTCCCTCGAGACAATGCCGCTGCGCATCG CACTTATC and GGTACCAGATCTTAGACTTCCCCTGAAGTAAATGG were 15 used to amplify ORF 7, and generate plasmid pCGN7771. The primers GAATTCGTCG ACACAATGTCATTACCAGACAATGCTTCT and TCTAGAGTCGACTTATAC AGATTCTCGATGCTGATAG were used to amplify ORF 8, and generate plasmid pCGN7775. The primers GAATTCGTCGACACAATGAATCCTACAGCAA CTAACGAA and TCTAGAGGATCCTTAGGCCATTCTTGGTTGGCTC were 20 used to amplify ORF 9, and generate plasmid pCGN7773.

The integrity of the PCR products was verified by DNA sequencing of the inserts of pCGN7771, PCGN8520, and pCGN7773. ORF 6 and ORF 8 were quite large in size. In order to avoid sequencing the entire clones, the center portions of the ORFs were replaced with restriction fragments of pEPA. The 6.6 kilobase *Pac*I/*Bam*HI fragment of pEPA containing the central portion of ORF 6 was ligated into *Pac*I/*Bam*HI digested pCGN7776 to yield pCGN7776B4. The 4.4 kilobase *Bam*HI/*Bgl*II fragment of pEPA containing the central portion of ORF 8 was ligated into *Bam*HI/*Bgl*II digested pCGN7775 to yield pCGN7775A. The regions flanking the pEPA fragment and the cloning junctions were verified by DNA sequencing.

Plasmid pCGN7771 was cut with *Xho*I and *Bgl*II and ligated to pCGN7770 after digestion with *Sal*I and *Bgl*II. The resultant napin/ORF 7 gene fusion plasmid was designated pCGN7783. Plasmid pCGN8520 was cut with *Xho*I and *Bgl*II and ligated to pCGN7770 after digestion with *Sal*I and *Bgl*II. The resultant napin/ORF 3 gene fusion plasmid was designated pCGN8528. Plasmid pCGN7773 was cut with *Sal*I and *Bam*HI and ligated to pCGN7770 after digestion with *Sal*I and *Bgl*II. The resultant napin/ORF 9 gene fusion plasmid was designated pCGN7785. Plasmid pCGN7775A was cut with *Sal*I and ligated to pCGN7770 after digestion with *Sal*I. The resultant napin/ORF 8 gene fusion plasmid was designated pCGN7782. Plasmid pCGN7776B4 was cut with *Xho*I and ligated to pCGN7770 after digestion with *Sal*I. The resultant napin/ORF 6 gene fusion plasmid was designated pCGN7786B4.

A binary vector for plant transformation, pCGN5139, was constructed from pCGN1558 (McBride and Summerfelt (1990) *Plant Molecular Biology*, 14:269-276). The polylinker of pCGN1558 was replaced as a *Hind*III/*Asp*718 fragment with a polylinker containing unique restriction endonuclease sites, *Ascl*, *Pac*I, *Xba*I, *Swa*I, *Bam*HI, and *Not*I. The *Asp*718 and *Hind*III restriction endonuclease sites are retained in pCGN5139. PCGN5139 was digested with *Not*I and ligated with *Not*I digested pCGN7786B4. The resultant binary vector containing the napin/ORF 6 gene fusion was designated pCGN8533. Plasmid pCGN8533 was digested with *Sse*8387I and ligated with *Sse*8387I digested pCGN7782. The resultant binary vector containing the napin/ORF 6 gene fusion and the napin/ORF 8 gene fusion was designated pCGN8535 (Figure 18).

The plant binary transformation vector, pCGN5139, was digested with *Asp*718 and ligated with *Asp*718 digested pCGN8528. The resultant binary vector containing the napin/ORF 3 gene fusion was designated pCGN8532. Plasmid pCGN8532 was digested with *Not*I and ligated with *Not*I digested pCGN7783. The resultant binary vector containing the napin/ORF 3 gene fusion and the napin/ORF 7 gene fusion was designated pCGN8534. Plasmid pCGN8534 was digested with *Sse*8387I and ligated with *Sse*8387I digested pCGN7785. The resultant binary vector containing the napin/ORF 3 gene fusion, the napin/ORF 7 gene fusion and the napin/ORF 9 gene fusion was designated pCGN8537 (Figure 19).

Vibrio constructs

The *Vibrio* ORFs for plant expression were all obtained using *Vibrio* cosmid #9 as a starting molecule. *Vibrio* cosmid #9 was one of the cosmids isolated from the *Vibrio* cosmid library using the *Vibrio* ORF 6 PCR product described in Example 1.

5 A gene encoding *Vibrio* ORF 7 (Figure 6) was mutagenized to introduce a *Sal*II site upstream of the open reading frame and *Bam*HI site downstream of the open reading frame using the PCR primers: TCTAGAGTCGACACAATGGCGGAATTAGCTG TTATTGGT and GTCGACGGATCCCTATTGTTCGTGGCTATG. A gene encoding *Vibrio* ORF 9 (Figure 6) was mutagenized to introduce a *Bam*HI site upstream 10 of the open reading frame and an *Xho*HI site downstream of the open reading frame using the PCR primers: GTCGACGGATCCACAATGAATATAGTAAGTAATCATTCGGCA and GTCGACCTCGAGTTAACACTCGTACGATAACTTGCC. The restriction sites were introduced using PCR, and the integrity of the mutagenized plasmids was verified 15 by DNA sequence. The *Vibrio* ORF 7 gene was cloned as a *Sal*-*Bam*HI fragment into the napin cassette of *Sal*-*Bgl*II digested pCGN7770 (Figure 17) to yield pCGN8539. The *Vibrio* ORF 9 gene was cloned as a *Sal*-*Bam*HI fragment into the napin cassette of *Sal*-*Bal*II digested pCGN7770 (Figure 17) to yield pCGN8543.

Genes encoding the *Vibrio* ORF 6 and ORF 8 were mutagenized to introduce *Sal*II sites flanking the open reading frames. The *Sal*II sites flanking ORF 6 were introduced 20 using PCR. The primers used were: CCCGGGTCGACACAATGGCTAAAAAGAACCA CCACATCGA and CCCGGGTCGACTCATGACATATCGTCAAAATGTCAGTGA. The central 7.3 kb *Bam*HI-*Xho*I fragment of the PCR product was replaced with the corresponding fragment from *Vibrio* cosmid #9. The mutagenized ORF 6 were cloned 25 into the *Sal*II site of the napin cassette of pCGN7770 to yield plasmid pCGN8554.

25 The mutagenesis of ORF 8 used a different strategy. A *Bam*HI fragment containing ORF 8 was subcloned into plasmid pHC79 to yield cosmid #9". A *Sal*II site upstream of the coding region was introduced on and adapter comprised of the oligonucleotides TCGACATGGAAAATTGCAGTAGTAGGTATTGCTAATT GTTC and CCGGGAACAAATTAGCAATACCTACTGCAATATTTCCATG. 30 The adapter was ligated to cosmid #9" after digestion with *Sal*II and *Xma*I. A *Sal*II site was introduced downstream of the stop codon by using PCR for mutagenesis. A DNA fragment containing the stop codon was generated using cosmid #9" as a template with the primers TCAGATGAACCTTATCGATAC and TCATGAGACGTCGACTTA

CGCTTCAACAATACT. The PCR product was digested with the restriction endonucleases *Clal* and *Aat*II and was cloned into the cosmid 9" derivative digested with the same enzymes to yield plasmid 8P3. The *Sal*I fragment from 8P3 was cloned into *Sal*I digested pCGN7770 to yield pCGN8515.

5 PCGN8532, a binary plant transformation vector that contains a *Shewannella* ORF 3 under control of the napin promoter was digested with *Not*I, and a *Not*I fragment of pCGN8539 containing a napin *Vibrio* ORF 7 gene fusion was inserted to yield pCGN8552. Plasmid pCGN8556 (Figure 23), which contains *Shewannella* ORF 3, and *Vibrio* ORFs 7 and 9 under control of the napin promoter was constructed by cloning the 10 *Sse8357* fragment from pCGN8543 into *Sse8387* digested pCGN8552.

15 The *Not*I digested napin/ORF 8 gene from plasmid pCGN8515 was cloned into a *Not*I digested plant binary transformation vector pCGN5139 to yield pCGN8548. The *Sse8387* digested napin/ORF 6 gene from pCGN8554 was subsequently cloned into the *Sse8387* site of pCGN8566. The resultant binary vector containing the napin/ORF 6 gene fusion and napin/ORF 8 gene fusion was designated pCGN8560 (Figure 22).

Example 5

Plant Transformation and PUFA Production

EPA production

20 The *Shewannella* constructs pCGN8535 and pCGN8537 can be transformed into the same or separate plants. If separate plants are used, the transgenic plants can be crossed resulting in heterozygous seed which contains both constructs.

pCGN8535 and pCGN8537 are separately transformed into *Brassica napus*. Plants are selected on media containing kanamycin and transformation by full length 25 inserts of the constructs is verified by Southern analysis. Immature seeds also can be tested for protein expression of the enzyme encoded by ORFs 3, 6, 7, 8, or 9 using western analysis, in which case, the best expressing pCGN8535 and pCGN8537 T1 transformed plants are chosen and are grown out for further experimentation and crossing. Alternatively, the T1 transformed plants showing insertion by Southern are crossed to one 30 another producing T2 seed which has both insertions. In this seed, half seeds may be analyzed directly from expression of EPA in the fatty acid fraction. Remaining half-seed

of events with the best EPA production are grown out and developed through conventional breeding techniques to provide *Brassica* lines for production of EPA.

Plasmids pCGN7792 and pCGN7795 also are simultaneously introduced into *Brassica napus* host cells. A standard transformation protocol is used (see for example 5 USPN 5,463,174 and USPN 5,750,871, however *Agrobacterium* containing both plasmids are mixed together and incubated with *Brassica* cotyledons during the cocultivation step. Many of the resultant plants are transformed with both plasmids.

DHA production

10 A plant is transformed for production of DHA by introducing pCGN8556 and pCGN8560, either into separate plants or simultaneously into the same plants as described for EPA production.

15 Alternatively, the *Shewanella* ORFs can be used in a concerted fashion with ORFs 6 and 8 of *Vibrio*, such as by transforming with a plant the constructs pCGN8560 and pCGN7795, allowing expression of the corresponding ORFs in a plant cell. This combination provides a PKS-like gene arrangement comprising ORFs 3, 7 and 9 of *Shewanella*, with an ORF 6 derived from *Vibrio* and also an ORF 8 derived from *Vibrio*. As described above, ORF 8 is the PKS-like gene which controls the identity of the final PUFA product. Thus, the resulting transformed plants produce DHA in plant oil.

20

Example 6

Transgenic plants containing the *Shewanella* PUFA genes

Brassica plants

25 Fifty-two plants cotransformed with plasmids pCGN8535 and pCGN8537 were analyzed using PCR to determine if the *Shewanella* ORFs were present in the transgenic plants. Forty-one plants contained plasmid pCGN8537, and thirty-five plants contained pCGN8535. 11 of the plants contained all five ORFs required for the synthesis of EPA. Several plants contained genes from both of the binary plasmids but appeared to be missing at least one of the ORFs. Analysis is currently being performed on approximately 30 twenty additional plants.

Twenty-three plants transformed with pCGN8535 alone were analyzed using PCR to determine if the *Shewanella* ORFs were present in the transgenic plants. Thirteen of

these plants contained both *Shewanella* ORF 6 and *Shewanella* ORF 8. Six of the plants contained only one ORF.

Nineteen plants transformed with pCGN8537 were alone analyzed using PCR to determine if the *Shewanella* ORFs were present in the transgenic plants. Eighteen of the 5 plants contained *Shewanella* ORF 3, *Shewanella* ORF 7, and *Shewanella* ORF 9. One plant contained *Shewanella* ORFs 3 and 7.

Arabidopsis

More than 40 transgenic *Arabidopsis* plants cotransformed with plasmids pCGN8535 and pCGN8537 are growing in our growth chambers. PCR analysis to 10 determine which of the ORFs are present in the plants is currently underway.

By the present invention PKS-like genes from various organisms can now be used to transform plant cells and modify the fatty acid compositions of plant cell membranes or plant seed oils through the biosynthesis of PUFAs in the transformed plant cells. Due to 15 the nature of the PKS-like systems, fatty acid end-products produced in the plant cells can be selected or designed to contain a number of specific chemical structures. For example, the fatty acids can comprise the following variants: Variations in the numbers of keto or hydroxyl groups at various positions along the carbon chain; variations in the numbers and types (*cis* or *trans*) of double bonds; variations in the numbers and types of branches 20 off of the linear carbon chain (methyl, ethyl, or longer branched moieties); and variations in saturated carbons. In addition, the particular length of the end-product fatty acid can be controlled by the particular PKS-like genes utilized.

All publications and patent applications mentioned in this specification are 25 indicative of the level of skill of those skilled in the art to which this invention pertains. All publications and patent applications are herein incorporated by reference to the same extent as if each individual publication or patent application was specifically and individually indicated to be incorporated by reference.

30 The invention now being fully described, it will be apparent to one of ordinary skill in the art that many changes and modifications can be made thereto without departing from the spirit or scope of the appended claims.

What is claimed is:

1. An isolated nucleic acid comprising:

a *Vibrio marinus* nucleotide sequence selected from the group consisting of the ORF 6, ORF 7, ORF 8 and ORF 9 as shown in Figure 6.

5

2. An isolated nucleic acid comprising:

a nucleotide sequence which encodes a polypeptide of a polyketide-like synthesis system, wherein said system produces a docosahexenoic acid when expressed in a host cell.

10 3. The isolated nucleic acid according to Claim 2, wherein said nucleotide sequence is derived from a marine bacterium.

4. The isolated nucleic acid according to Claim 2, wherein said nucleotide sequence is a *Vibrio marinus* ORF 8 as shown in Figure 6.

15

5. An isolated nucleic acid comprising:

a nucleotide sequence which is substantially identical to a sequence of at least 50 nucleotides of a *Vibrio marinus* nucleotide sequence selected from the group consisting of ORF 6, ORF 7, ORF 8 and ORF 9 as shown in Figure 6.

20

6. A recombinant microbial cell comprising at least one copy of an isolated nucleic acid according to Claim 1 or Claim 2.

25 7. The recombinant microbial cell according to Claim 6, wherein said cell comprises each element of a polyketide-like synthesis system required to produce a long chain polyunsaturated fatty acid.

8. The recombinant microbial cell according to Claim 7, wherein said cell is a eukaryotic cell.

30

9. The recombinant microbial cell according to Claim 8, wherein said eukaryotic cell is a fungal cell, an algae cell or an animal cell.

10. The recombinant microbial cell according to Claim 9, wherein said fungal cell is a yeast cell and said algae cell is a marine algae cell.

5 11. The recombinant microbial cell according to Claim 6, wherein said cell is a prokaryotic cell.

12. The recombinant microbial cell according to Claim 11, wherein said cell is a bacterial cell or a cyanobacterial cell.

10 13. The microbial cell according to Claim 6, wherein said recombinant microbial cell is enriched for 22:6 fatty acids as compared to a non-recombinant microbial cell which is devoid of said isolated nucleic acid.

15 14. A method for production of docosahexenoic acid in a microbial cell culture, said method comprising:

20 growing a microbial cell culture having a plurality of microbial cells, wherein said microbial cells or ancestors of said microbial cells were transformed with a vector comprising one or more nucleic acids having a nucleotide sequence which encodes a polypeptide of a polyketide synthesizing system, wherein said one or more nucleic acids are operably linked to a promoter, under conditions whereby said one or more nucleic acids are expressed and docosahexenoic acid is produced in said microbial cell culture.

15. A method for production of a long chain polyunsaturated fatty acid in a plant cell, said method comprising:

25 growing a plant having a plurality of plant cells, wherein said plant cells or ancestors of said plant cells were transformed with a vector comprising one or more nucleic acids having a nucleotide sequence which encodes one or more polypeptides of a polyketide synthesizing system which produces a long chain polyunsaturated fatty acid, wherein each of said nucleic acids are operably linked to a promoter functional in a plant cell, under conditions whereby said polypeptides are expressed and a long chain polyunsaturated fatty acid is produced in said plant cells.

16. The method according to Claim 15, wherein said long chain polyunsaturated fatty acid produced in said plant cells is a 20:5 and 22:6 fatty acid.

17. The method according to Claim 15, wherein said nucleic acids comprise 5 nucleotide sequences encoding any one of the polypeptides selected from the group consisting of *Vibrio marinus* ORF 6, ORF 7, ORF 8 and ORF 9 as shown in Figure 6 and *Shewanella putrefaciens* ORF 3, ORF 6, ORF 7, ORF 8 and ORF 9 as shown in Figure 4.

18. The method according to Claim 15, wherein said nucleic acid constructs are derived 10 from two or more polyketide synthesizing systems.

19. A recombinant plant cell which produces an long chain polyunsaturated fatty acid exogenous to said plant cell, wherein said recombinant plant cell is produced according to a method comprising:

15 transforming a plant cell or an ancestor or said plant cell with a vector comprising one or more nucleic acids having a nucleotide sequence which encodes one or more polypeptides of a polyketide synthesizing system which produces a long chain polyunsaturated fatty acid, wherein each of said nucleic acids are operably linked to a promoter functional in said plant cell whereby a recombinant plant cell is obtained; and 20 growing said recombinant plant cell under conditions whereby said polypeptides are expressed and a long chain polyunsaturated fatty acid is produced in said plant cell.

20. The recombinant plant cell according to Claim 19, wherein said recombinant plant cell is a recombinant seed cell.

25 21. The recombinant plant cell according to Claim 20, wherein said recombinant seed cell is a recombinant embryo cell.

30 22. The method according to Claim 15, wherein said long chain polyunsaturated fatty acid produced in said plant cells is eicosapentenoic acid.

23. The method according to Claim 15, wherein said long chain polyunsaturated fatty acid produced in said plant cells is docosahexenoic acid.

24. The recombinant plant cell according to Claim 19, wherein said recombinant plant cell is from a plant selected from the group consisting of *Brassica*, soybean, safflower, and sunflower.

5 25. A plant oil produced by a recombinant plant cell according to Claim 19, wherein said plant oil comprises eicosapentenoic acid.

26. A plant oil produced by a recombinant plant cell according to Claim 19, wherein said plant oil comprises docosahexenoic acid.

10

27. The plant oil according to Claim 25 or Claim 26, wherein said plant oil is encapsulated.

28. A dietary supplement comprising a plant oil according to Claim 27.

15

29. A recombinant *E. coli* cell which produces docosahexenoic acid.

30. A plant oil comprising eicosapentenoic acid.

31. A plant oil comprising docosahexenoic acid.

20

32. The recombinant microbial cell according to Claim 12, wherein said bacterial cell is a lactobacillus cell.

Fig. 1 Organization of *Shewanella EPA* Genes and Clones Obtained from the Sagami Chemical Institute.

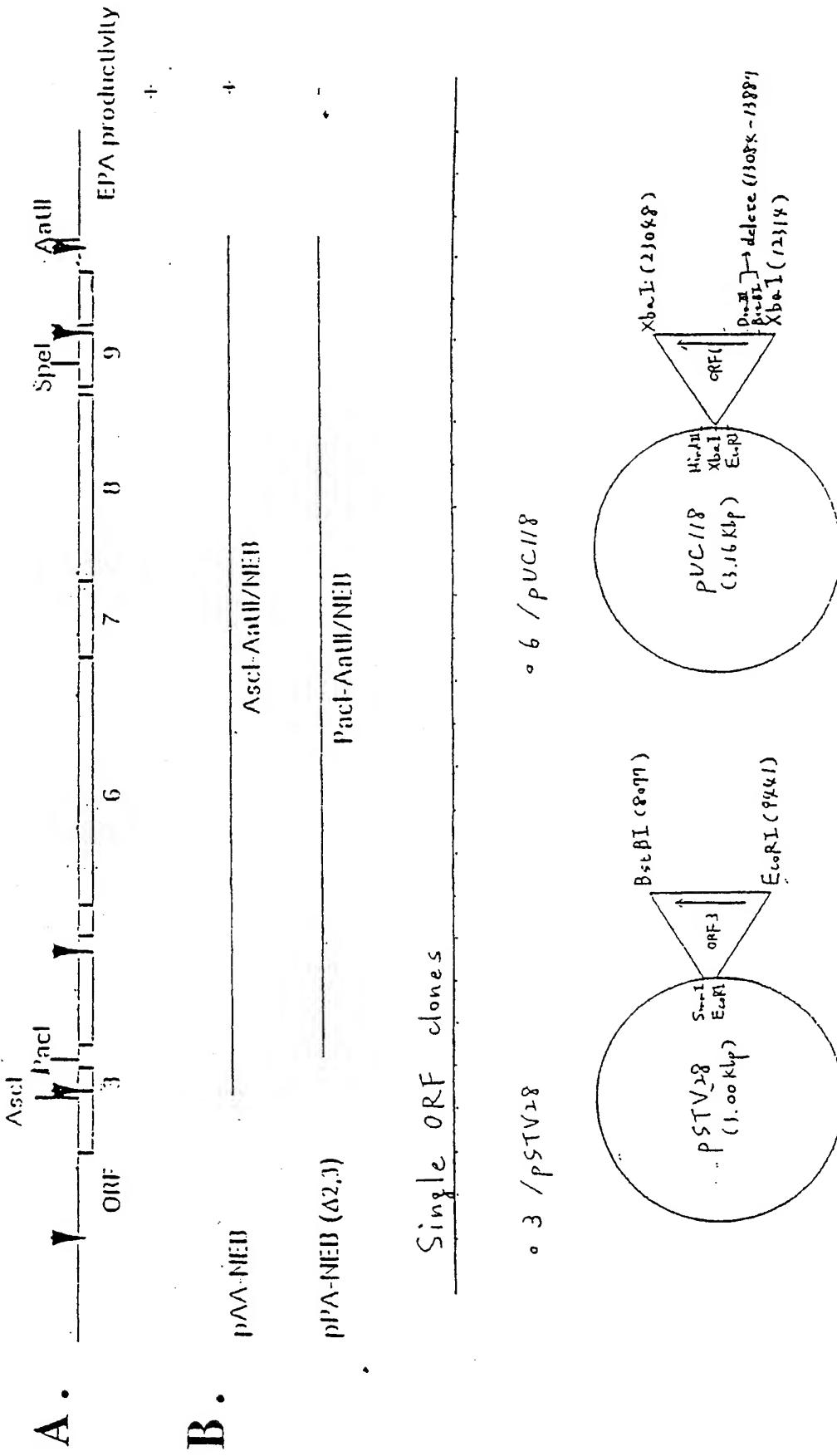
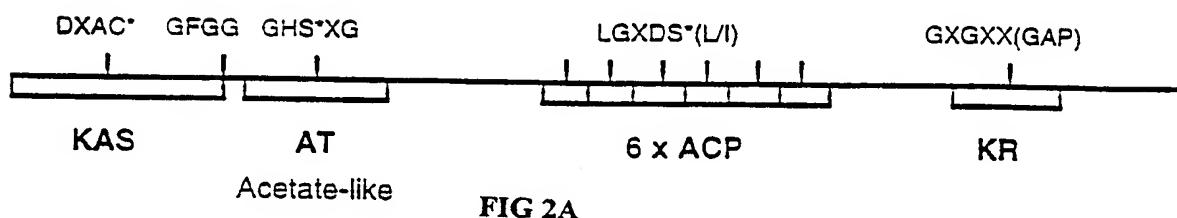


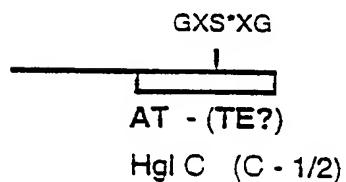
Fig. 2

SHIWANELLA EPA C RFs**Motifs - Domains - Homologies**

Orf6 8.3 KB - 293 kD



Orf7 2.3 KB - 84 kD

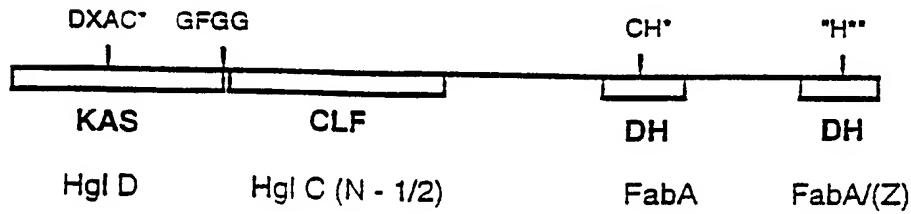


Orf3 0.8 KB - 30 kD

Het I - pantetheine transferase

FIG 2E

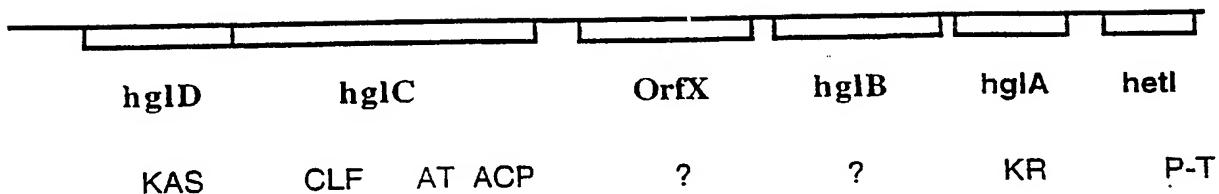
Orf8 6.0 KB - 217 kD



Orf9 1.6 KB - 59 kD

Anabeana - OrfX homolog

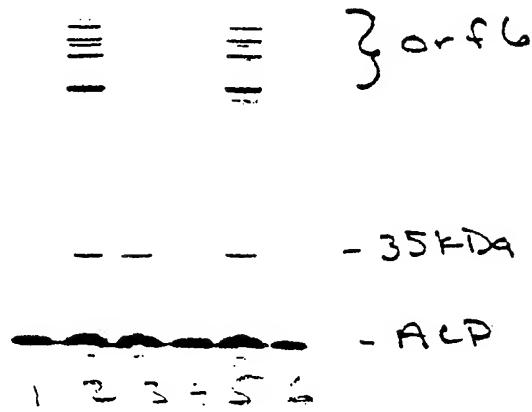
FIG 2D



Anabeana "PKS" Genes Involved in Heterocyst Glycolipid Synthesis**

Fig 3. Orf3 Encodes a Phosphopantetheine Transferase

1. pUC19
3. pAA-Neb (EPA +)
2. pPA-NEB (Δ Orf3)
4. Orf6 subclone
5. Orf6 + Orf3 subclones
6. Orf3 subclone



Autoradiograph of [C^{14}] β -Alanine labelled proteins from *E. coli* (strain SJ16) cells transformed with the above listed plasmids. Cells were grown in the presence of [C^{14}] β -alanine and the appropriate antibiotics. Proteins were extracted, separated by SDS-PAGE and transferred to a PVDF membrane prior to autoradiography. ACP and an unknown (but previously observed) 35 kD protein were labelled in all of the samples. The high molecular mass proteins detected in lanes 2 and 5 are full-length (largest band) and truncated products of the *Shewanella* Orf6 gene (confirmed by Western analysis - data not shown). *E. coli* strain SJ16 is conditionally blocked in β -alanine synthesis.

Sequence Range: 1 to 37895

20	40	60	80
GATCTCTTAC AAAGAAAAT TCTCAATGTG AATTTAACCT TAATTCGGT TAATTACGGC CTGATAGAGC ATCACCCAAT			
100	120	140	160
CAGCCATAAA ACTGTAAAGT GGGTACTCAA AGGTGGCTGG GCGATTCTTC TCAAATACAA AGTGCCCAAC CCAAGCAAAT			
180	200	220	240
CCATATCCGA TAACAGGTAA AAGTAGCAAAT AAACCCCAGC GCTGAGTTAG TAATACATAA GCGAATAATAA GGATCACTAA			
260	280	300	320
ACTACTGCCG AAATAGTGTAA ATATTCGACA GTTTCTATGC TGATGTTGAG ATAATAAAA AGGGTAAAAT TCAGCAAAAG			
340	360	380	400
AACGATAGCG CTTACTCATT ACTCACACCT CGGTAAAAAA GCAACTCGCC ATTAACTTGG CCAATCGTCA GTTGTCTAT			
420	440	460	480
CGTCTCAAAG TTATGCCGAC TAAATAACTC TATATGTGCA TTATGATTAG CAAAAACTCC GATACCATCA AGATGAAGTT			
500	520	540	560
GTTCATCACA CCAACTCAAA ACTGCGTCGA TAAGCTTACT CCCATAGCCC TTGCCTTGCT CCACATTGC GATAGCAATA			
580	600	620	640
AACTGTAAAAA TCCCACATTG CCCACTTGGT AAGCTCTCTA TAATCTGATT TTCTTTGTTA ATAAGTGCCT GAGTTGAATA			
660	680	700	720
CCAACCAGTA CTTAACARCA TCTTTAAACG CCAATGCCAA AAACGGCGTT CACCTAAGGG AACCTGCTGA GTCACTATGC			
740	760	780	800
AGGCTACGCC TATCAATCTA TCCCCAACGA ACATACCAAT AAGTGCTTGC TCCTGTTGCC AGAGCTCATT GAGTTCTTCT			
820	840	860	880
CGAATAGCCC CGCGAAGCTT TTGCTCATAAC TGCCTTGTG CACCACTAAA AAGTGTTCG ATAAAAAAGG GATCATCATG			
900	920	940	960
ATAGGCGTTA TAGAGAATAG AGGCTGCTAT GCGTAAATCT TCTGCCGTGA GATAAACTGC ACGACACTCT TCCATGGCTT			
980	1000	1020	1040
GATCTTCCAT TGTTATTGTC CTTGACCTTG ATCACACAAAC ACCAATGTAA CAAGACTGTA TAGAAGTGCA ATTAATAATC			
1060	1080	1100	1120
AATTCGTGCA TAAAGCAGST CAGCATTCT TTGCTAAACAA AGCTTTATTG GCTTTGACAA AACTTTGCCT AGACTTTAAC			
1140	1160	1180	1200
GATAGAAATC ATAATGAAAG AGAAAAGCTA CAACCTAGAG GGGAAATAATC AAACAACCTGC TAAGATCTAG ATAATGTAAT			
1220	1240	1260	1280
AAACACCGAG TTTATCGACC ATACTTAGAT AGAGTCATAG CAACGAGAAT AGTTATGGAT ACAACGCCCG AAGATCTATC			
1300	1320	1340	1360
ACACCTGTTT TTACAGCTAG GATTAGCAAA TGATCAACCC GCAATTGAAC AGTTTATCAA TGACCATCAA TTAGCGGACA			
1380	1400	1420	1440
ATATATTGCT ACATCAAGCA AGCTTTGGAA GCCCATCGCA AAAGCACTTC TTAATTGAGT CATTAAATGA AGATGCCAG			
1460	1480	1500	1520
TGGACCGAAG TCATCGACCA CTTAGACACC TTATTAAGAA AAAACTAACCC ATTACAACAG CAACTTTAAA TTTGCCGTA			
1540	1560	1580	1600
AGCCATCTTC CCCCCACCCCA CAACAGCGTT GTTGCTTATG ACCACTGGAG TACATTCCCTC TTTAGTCGTT TTACCATCAC			
1620	1640	1660	1680
CATGGGTACG TTGAGTGCAGA TAAAAAAGCA CATAAACTTC TTTATCGGCC TGAATATAGG CTTCGTTAAA ATCAGCTGTT			
1700	1720	1740	1760
CCCATTAAAG TAACCACTTG CTCTTTACTC ATGCCTAGAG ATATCTTGT CAAATTGTCA CGGTTTTAT CTTGAGTTT			

Fig. 4
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1780 1800 1820 1840
 * * * *
 CTCCCAAGCA CCGTGATTAT CCCAGTCAGA TTCCCCATCA CCAACATTGA CCACACAGCC CGTTAGCCCT AAGCTTGCAA
 1860 1880 1900 1920
 * * * *
 TCCCCAAACACA TGCTAAACCT AATAATTAT TTTTCATTT AACTTC~~T~~GT TATGACATTA TTTTGCTTA GAAGAAAAGC
 1940 1960 1980 2000
 * * * *
 AACTTACATG CCAAAACACA AGCTGTTGTT TTAAATGACT TTATTTATTA TTAGCCTTT AGGATATGCC TAGAGCAATA
 2020 2040 2060 2080
 * * * *
 ATAATTACCA ATGTTTAAGG AATTGACTA ACTATGAGTC CGATTGAGCA AGTGCTAACAA GCTGCTAAAA AAATCAATGA
 2100 2120 2140 2160
 * * * *
 ACAAGGTTAGA GAACCAACAT TAGCATTGAT TAAAACAAA CTTGGTAATA GCATCCCAAT GCGCGAGTTA ATCCAAGGTT
 2180 2200 2220 2240
 * * * *
 TCGAACAGTT TAAGTCTATG AGTGCAGAAG AAAGACAAGC AATACTTAGC AGCTTAGCAA CAGCAAAAGA AACTCAATAT
 2260 2280 2300 2320
 * * * *
 GGTCAATCAA GCTTATCTCA ATCTGAACAA GCTGATAGGA TCCTCCAGCT AGAAAACGCC CTCATGAAT TAAGAAACGA
 2340 2360 2380 2400
 * * * *
 ATTTAATGGG CTAAAAAGTC AATTGATAAA CTTACACAA AACCTGATGA ATAAAGAGCC TGACACCAAA TGCAATGTAAT
 2420 2440 2460 2480
 * * * *
 TGAACTACGA TTTGAATGTT TTGATAACAC CACGATTACT GCAGCAGAAA AAGCCATTAA TGTTTGCTT GAAGCTTATC
 2500 2520 2540 2560
 * * * *
 GAGCCAATGG CCAGGTTCTA GGTCGTGAAT TTGCCGTTGC ATTTAACGAT GGTCAGTTA AAGCACCGAT GTTAACCCCA
 2580 2600 2620 2640
 * * * *
 GAAAAAAAGCA GCTTATCTAA ACGCTTTAAT AGTCCTGGG TAAATAGTGC ACTCGAAGAG CTAACCGAAG CCAAATTGCT
 2660 2680 2700 2720
 * * * *
 TGCGCCACGT GAAAAGTATA TTGGCCAAGA TATTAATTCT GAAGCATCTA GCCAAGACAC ACCAAGTTGG CAGCTACTTT
 2740 2760 2780 2800
 * * * *
 ACACAAAGTTA TGTGCACATG TGTCACCAC TAAGAAATGG CGACACCTTG CAGCCTATTG CACTGTATCA AATTCCAGCA
 2820 2840 2860 2880
 * * * *
 ACTGCCAACG GCGATCATAA ACGAATGATC CGTTGGCAAA CAGAATGGCA AGCTTGTGAT GAATTGCAA TGGCCGCAGC
 2900 2920 2940 2960
 * * * *
 TACTAAAGCT GAATTGCGG CACTTGAAGA GCTAACCGAT CATCAGAGTG ATCTATTTAG GCGTGGTTGG GACTTACGTG
 2980 3000 3020 3040
 * * * *
 GCAGAGTCGA ATACTTGACG AAAATTCCGA CCTATTACTA TTTATACCGT GTTGGCGGTG AAAGCTTAGC AGTAGAAAAG
 3060 3080 3100 3120
 * * * *
 CAGCGCTCTT GTCCTAAGTG TGGCAGTCAA GAATGGCTGC TCGATAAACCC ATTATTGGAT ATGTTCCATT TTCGCTGTGA
 3140 3160 3180 3200
 * * * *
 CACCTGCCGC ATCGTATCTA ATATCTCTTG GGACCATTAA TAACTCTTCC GAGTCTTATC ACACTAGAGT TTAGTCAGCA
 3220 3240 3260 3280
 * * * *
 TAAAAATGGC GCTTATATTT CAATTAAAAG AAATATAAGC GCCATTTCA TCGATACTAT ATATCAGCAG ACTATTTCC
 3300 3320 3340 3360
 * * * *
 GCGTAAATTAA GCCCACATTA ATTTCATTCT TTGCCAGATC CCTGGATGAT CTAGTTGTGG CATCGACTCT TCAATAGGTT
 3380 3400 3420 3440
 * * * *
 TAACCGCAGG TGTAACCCCTT GGAGTCAATT CGTTTATAAA CTCGTTAAA CTGTCACTTA ATTTAACGCT TTGTACTTCA
 3460 3480 3500 3520
 * * * *
 CCTGGAATTAA CAATCCATAC GCTGCCATCA CTATTATTAA CCGTCAACAT TTTATCTTCA TCATCAAGAA TACCAATAAA

Fig. 4
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3540	3560	3580	3600				
*	*	*	*				
CCAAGTCGGC	TCTTGCTTAA	GCTTTCTCTT	CATCATTAAA	TGACCAATGA	TGTTTGTG	TAAGTATTCA	AAATCAGTT
3620	3640	3660	3680				
*	*	*	*				
GATCCCACAC	TTGGATTAGC	TCACCTTGGC	CCCATTGTGA	GTCAAAAAAT	AGCGGTGCAG	AAAAATGACT	GCCAAAAAAAT
3700	3720	3740	3760				
*	*	*	*				
GGATTAATTT	CTGCAGATAA	TGTCATTCA	AGTGCTGTT	CAACATTAGC	AAATTCAACCA	GGTTGTTGAC	GTACAACCGA
3780	3800	3820	3840				
*	*	*	*				
TTGCCAAAAC	ACTGCGCCAT	CGGAGCCCC	TTCGGCAGA	ACACACTCAG	ACTTTGTCC	TTGGCATAA	TATCTTGGCT
3860	3880	3900	3920				
*	*	*	*				
GTTCACCAAG	CTTATCCATG	TAGGCTTGTG	GATATTAGA	AAAAAAAGA	TCTAAAGCG	GTAAAGAAGA	CACTTAAGCC
3940	3960	3980	4000				
*	*	*	*				
AGTTCCAAAAA	TCAGTTATAA	TAGGGTCTA	TTTGACATG	GAAACCGTAT	TGATGACACA	ACATCATGAT	CCCTACAGTA
4020	4040	4060	4080				
*	*	*	*				
ACGCCCGA	ACTTTCTGAA	TTAACCTTAG	GAAAGTCGAC	CGGTTATCAA	GAGCAGTATG	ATGCATCTT	ACTACAAGCG
4100	4120	4140	4160				
*	*	*	*				
TGCCCGTAA	ATTAAACCGT	GATGCTATCG	GTCTAACCAA	TGAGCTACCT	TTTCATGGCT	GTGATATTG	GACTGGCTAC
4180	4200	4220	4240				
*	*	*	*				
GAACTGTCTT	GGCTAAATGC	TAAAGGCAAG	CCAATGATTG	CTATTGCAGA	CTTTAACCTA	AGTTTTGATA	GTAAAAATCT
4260	4280	4300	4320				
*	*	*	*				
GATCGAGTCT	AACTCGTTA	AGCTGTATTT	AAACAGCTAT	AACCAAACAC	GATTGATAG	CGTTCAAGCG	GTTCAAGAAC
4340	4360	4380	4400				
*	*	*	*				
GTTTAACTGA	AGACTTAAGC	GCCTGTCCCC	AAGGCACAGT	TACGGTAAAA	GTGATTGAAC	CTAAGCAATT	TAACCACCTG
4420	4440	4460	4480				
*	*	*	*				
AGAGTGGTTG	ATATGCCAGG	TACCTGCATT	GACGATTAG	ATATTGAAGT	TGATGACTAT	AGCTTTAACT	CTGACTATCT
4500	4520	4540	4560				
*	*	*	*				
CACCGACAGT	GTTGATGACA	AACTCATGGT	TGCTGAAACG	CTAACGTCAA	ACTTATTGAA	ATCAAACATGC	CTAATCACTT
4580	4600	4620	4640				
*	*	*	*				
CTCAGCCTGA	CTGGGGTACA	GTGATGATCC	GTTATCAAGG	GCCTAAGATA	GACCGTAAA	AGCTACTTAG	ATATCTGATT
4660	4680	4700	4720				
*	*	*	*				
TCATTTAGAC	AGCACAAATGA	ATTTCATGAG	CAGTGTGTTG	AGCGTATATT	TGTTGATTAA	AAGCACTATT	GCCAAATGTGC
4740	4760	4780	4800				
*	*	*	*				
CAAACCTACT	GTCTATGCAC	GTTATACCG	CCGTGGTGGT	TTAGATATCA	ACCCATATCG	TAGCGACTTT	GAAAACCTG
4820	4840	4860	4880				
*	*	*	*				
CAGAAAATCA	GCGCCTAGCG	AGACAGTAAT	TGATTGAGT	ACCTACAAA	AACAATGCCT	ATAAGCCAAG	CTTATGGGCA
4900	4920	4940	4960				
*	*	*	*				
TTTTTATATT	ATCAACTTGT	CATCAAACCT	CAGCCCCAA	GCCTTTAGT	TTTATCGCTA	ATTAAGCCG	CTCTCTCAGC
4980	5000	5020	5040				
*	*	*	*				
CAAATATTG	CAGGATTGG	CTGTAATTAA	TGGCTCCACA	CCATGAAATA	CTCTATCGGC	TCTACCGCAA	AAGGTAAGTC
5060	5080	5100	5120				
*	*	*	*				
AAATACCTGT	AAGCCAAACA	GCTTGGCATA	TTCGTCAGTG	TGGGCTTTG	ACCGCGATAGC	TAACGCATCA	CTTTTGAGG
5140	5160	5180	5200				
*	*	*	*				
CAACCGACAT	CATACTTAAT	ATTGATGATT	GCTCGCTGTG	CATTGCTT	GCCGGTAACA	CCTGTTAGT	CAGCAAGTCC
5220	5240	5260	5280				
*	*	*	*				
GCAACACTTA	AATTGTAGCG	GCGCATCTTA	AAAATAATAT	GCTTTTCATT	AAAGTATTGC	TCTTGCCTCA	ACCCACCTTG
5300	5320	5340	5360				

Fig 4
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GATCCTGGG TGAGCATTG GTGCCACACA AACTAATTG TCCGCATTA CTTTTGACT CTTAAATGCC GCAGATTCTG
 5380 5400 5420 5440
 * * * *
 GCAGCCAAAT ATCTAAGGCT AAATCCACCT TTTCTAGTTG TAGGTCCATC TGCAACTCTT CTTCAATGAG CGGGGGCTCA
 5460 5480 5500 5520
 * * * *
 CGAAATACAA TATTAATTGC AGTGCCTGT AACACTTGCT CAATTTGATC TTGCAAGAGT TGTATTGCCG ACTCGCTGGC
 5540 5560 5580 5600
 * * * *
 ATACACATAA AAAGTTCGCT CACTTGAAGT GGGGTCAAAT GCTTCAAAGC TAGTCGAAC TTGCTCAATT GTTGACATAG
 5620 5640 5660 5680
 * * * *
 CGCCCGCAG CTGTTGATAA AGCGTCATCG CACTTGCCTG AGGTTAACT CCCCTACCCA CTCGAGTAA CAACTCTTCT
 5700 5720 5740 5760
 * * * *
 CCAACAAATAC TTTTAGCCT CGAAATCGCA TTACTAACCG ACGACTGAGT CAAATCCAGC TCTTCTGCCG CCCGGCTAAA
 5780 5800 5820 5840
 * * * *
 AGATGAGGTG CGATACACCG CAGTAAAAAC GCGAAATAAA TTAAGATCAA AAGCTTTTG CTGCGACATA AATCAGCTAT
 5860 5880 5900 5920
 * * * *
 CTCCCTATCC TTATCCTTAT CCTTATAAAA AGTTAGCTCC AGAGCACTCT AGCTAAAAAA CAACTCAGCG TATTAAGCCA
 5940 5960 5980 6000
 * * * *
 ATATTTGGG AACTCAATTA ATATTCATAA TAAAAGTATT CATAATATAA ATACCAAGTC ATAATTAGC CCTAATTATT
 6020 6040 6060 6080
 * * * *
 AATCAATTCA AGTTACCTAT ACTGGCCTCA ATTAAGCAAA TGTCTCATCA GTCTCCCTGC AACTAAATGC AATATTGAGA
 6100 6120 6140
 * * * *
 CATAAAAGCTT TGAACGTATT CAATCTTACG AGCGTAACCTT ATG AAA CAG ACT CTA ATG GCT ATC TCA ATC ATG
 M K Q T L M A I S I M >
 6160 6180 6200
 * * * *
 TCG CTT TTT TCA TTC AAT GCG CTA GCA GCG CAA CAT GAA CAT GAC CAC ATC ACT GTT GAT TAC GAA
 S L F S F N A L A A Q H E H D H I T V D Y E >
 6220 6240 6260 6280
 * * * *
 GGG AAA GCC GCA ACA GAA CAC ACC ATA GCT CAC AAC CAA GCT GTA GCT AAA ACA CTT AAC TTT GCC
 G K A A T E H T I A H N Q A V A K T L N F A >
 6300 6320 6340
 * * * *
 GAC ACG CGT GCA TTT GAG CAA TCG TCT AAA AAT CTA GTC GCC AAG TTT GAT AAA GCA ACT GCC GAT
 D T R A F E Q S S K N L V A K F D K A T A D >
 6360 6380 6400
 * * * *
 ATA TTA CGT GCC GAA TTT GCT TTT ATT AGC GAT GAA ATC CCT GAC TCG GTT AAC CCG TCT CTC TAC
 I L R A E F A F I S D E I P D S V N P S L Y >
 6420 6440 6460 6480
 * * * *
 CGT CAG GCT CAG CTT AAT ATG GTG CCT AAT GGT CTG TAT AAA GTG AGC GAT GGC ATT TAC CAG GTC
 R Q A Q L N M V P N G L Y K V S D G I Y Q V >
 6500 6520 6540
 * * * *
 CGC GGT ACC GAC TTA TCT AAC CTT ACA CTT ATC CGC AGT GAT AAC GGT TGG ATA GCA TAC GAT GTT
 R G T D L S N L T L I R S D N G W I A Y D V >
 6560 6580 6600
 * * * *
 TTG TTA ACC AAA GAA GCA GCA AAA GCC TCA CTA CAA TTT GCG TTA AAG AAT CTA CCT AAA GAT GGC
 L L T K E A A K A S L Q F A L K N L P K D G >
 6620 6640 6660 6680
 * * * *
 GAT TTA CCC GTT GTT GCG ATG ATT TAC TCC CAT AGC CAT GCG GAC CAC TTT GGC GGA GCT CGC GGT
 D L P V V A M I Y S H S H A D H F G G A R G >
 6700 6720 6740
 * * * *
 GTT CAA GAG ATG TTC CCT GAT GTC AAA GTC TAC GGC TCA GAT AAC ATC ACT AAA GAA ATT GTC GAT
 V Q E M F P D V K V Y G S D N I T K E I V D >

Fig. 4
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Fig. 4
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7940 * * * * * 7960 * * * * * 7980 * * * * * 8000 * * * * *
 GTT AAT AAA GCT GAC GTT AAC CGC ATC TTA CTT GGC CAA GTA ACC CTA AAA GCG TTA TTA GCC AGC
 V N K A D V N R I L L G Q V T L K A L L A S >
 8020 * * * * * 8040 * * * * * 8060 * * * * *
 GGC GAT GCC AAG CTC ACT GGT GAT AAA ACG GCA TTT AGT AAA ATA GCC GAT AGC ATG GTC GAG TTT
 G D A K L T G D K T A F S K I A D S M V E F >
 8080 * * * * * 8100 * * * * * 8120 * * * * * 8140 * * * * *
 ACA CCT GAC TTC GAA ATC GTA CCA ACG CCT GTT AAA TGAGGCA TTAATCTCAA CAAGTGCAAG CTAGACATAA
 T P D F E I V P T P V K >
 8160 * * * * * 8180 * * * * * 8200 * * * * *
 AAATGGGGCG ATTAGACGCC CCATTTTTA TGCAATTTG AACTA GCT AGT CTT AGC TCA AGC TCG AAC AAC
 <S T K A S A R V V >
 8220 * * * * * 8240 * * * * * 8260 * * * * *
 AGC TTT AAA ATT CAC TTC TTC TGC TGC AAT ACT TAT TTG CTG ACA CTG ACC AAT ACT CAG TCC AAA
 <A K F N V E E A A I S I Q Q C Q G I S L A F >
 8280 * * * * * 8300 * * * * * 8320 * * * * * 8340 * * * * *
 ACG ATA ACT ATC ATC AAG ATG GCC CAG TAA ACA ATG CCA ATT ATC AGC AGC GTT CAT TTG CTG TTC
 <R Y S D D L H G L L C H W N D A A N M Q Q E >
 8360 * * * * * 8380 * * * * * 8400 * * * * *
 TTT AGC CTC AAT CAA ACC TAA ACC AGA CTT TTG TGG CTC AGC GTT AGG CTT ATT AGA ACT CGA CTC
 <K A E I L G L G S K Q P E A N P K N S S S E >
 8420 * * * * * 8440 * * * * * 8460 * * * * *
 TAG TAA AGC AAG ACC AAT ATC TTG TTT TAA CAA AAC CTG TCG CTG ATT AAG TTG ATG CTC AAC CTT
 <L L A L G I D Q K L L V Q R Q N L Q H E V K >
 8480 * * * * * 8500 * * * * * 8520 * * * * * 8540 * * * * *
 GTG ATC CGC AAT AGC ATC GGA AAT ATC AAC ACA ATG GCT CAA GCT TTT AGG TGC ATT AAC TCC AAG
 <H D A I A D S I D V C H S L S K P A N V G L >
 8560 * * * * * 8580 * * * * * 8600 * * * * *
 AAA AGT TTC GCT CAG TGC AGA GAA GTC AAA CGC AAA AGA TTT TAG CGA TAA TGC CAG CCC AAG TCC
 <F T E S L A S F D F A F S K L S L A L G L G >
 8620 * * * * * 8640 * * * * * 8660 * * * * *
 TTT CGC TTT AAT GTA AGA CTC CTT GAG CGC CCA CAA ATC AAA AAA GCG GTC TCG CTG CAA GGC CTC
 <K A K I Y S E K L A W L D F F R D R Q L A E >
 8680 * * * * * 8700 * * * * * 8720 * * * * * 8740 * * * * *
 TGG TAA CGC TAA CAA GGC TCG CTT TTC TGA TTC AGA GAA ATA ATG ACT AAG AAT AGA GTG GAT ATT
 <P L A L L A R K E S E S F Y H S L I S H I N >
 8760 * * * * * 8780 * * * * * 8800 * * * * *
 GGT GCT GTT ACG GCA ACG CTC AAT GTC GAC GCC AAA CTC AAT ACT AGC AGA GTC AGT TTC CTC CTT
 <T S N R C R E I D V G F E I S A S D T E E K >
 8820 * * * * * 8840 * * * * * 8860 * * * * *
 GCT TGC CTG ACT GGC GCC TTT ATT ATC AGC AGT GCA AAT GCC TAC TAA TAG CCA ATC TCC ACT ATG
 <S A Q S A G K N D A T C I G V L L W D G S H >
 8880 * * * * * 8900 * * * * * 8920 * * * * *
 ACT CAC ATT AAA GTG GAC CCC GGT TTG AGC AAA TTG CGC ATC ACT CAA TCT AGG CTT ACC TTT GTC
 <S V N F H V G T Q A F Q A D S L R P K G K D >
 8940 * * * * * 8960 * * * * * 8980 * * * * * 9000 * * * * *
 GCC ATA TTC AAA GCG CCA TTC ATT GGG GCG TAT TTC ACT ATG TTG TGA CAA TAA AGC GCG CAA ATA
 <G Y E F R W E N P R I E S H Q S L L A R L Y >
 9020 * * * * * 9040 * * * * * 9060 * * * * * 9080 * * * * *
 GCC TCT TAC CAT TAAA CCTTGAGTTT TAGCTCTTG TTTAATGTAG CGATTAACCT TAATTAACTC ATCTTCAGGC
 <G R V M >
 9100 * * * * * 9120 * * * * * 9140 * * * * * 9160 * * * * *

←
0.43
Fig. 4
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9180	9200	9220	9240
AGAAGTTAGC AATTTTCAG GACACTCTT AAAGCAACAA ACATAACCCC TATTTTACCC AATTTAAGAT CAAAACCTAAA			
9260	9280	9300	9320
GCCAAAACCA ATTGAGAATA GTGTCAAACT AGCTTTAAAG GAAAAAAATA TAAAAAGAAC ATTATACTTG TATAAATTAT			
9340	9360	9380	9400
TTTACACACC AAAGCCATGA TCTTCACAAA ATTAGCTCCC TCTCCCTAAACAAAGATTGA ATAAAAAAAT AAACCTTAAAC			
9420	9440	9460	9480
TTTCATATAG ATAAAACAAA CCAATGGGAT AAAGTATATT GAATTCAATT TTAAGGAAAA ATTCAAATTG AATTCAAGCT			
9500	9520	9540	9560
CTTCAGTAAA AGCATATTTT GCCGTTAGTG TGAAAAAAA CAAATTAAA AACCAACATA GAACAAATAA GCAGACAATA			
9580	9600	9620	9640
AAACCAAGGC GCAACACAAA CAACCGCGCTT ACAATTTCAC CAAAAAGCA ACAAGAGTAA CGTTTAGTAT TTGGATATGG			
9660	9680	9700	
TTATTGTAAT TGAGAATTT ATAACAATTA TATTAAGGGATG AGT ATG ATG TTT TTA AAT TCA AAA CTT TCG CGC	M S M F L N S K L S R >		
9720	9740	9760	
TCA GTC AAA CTT GCC ATA TCC GCA GGC TTA ACA GCC TCG CTA CCT ATG CCT GTT TTT GCA GAA GAA	S V K L A I S A G L T A S L A M P V F A E E >		
9780	9800	9820	9840
ACT GCT GCT GAA GAA CAA ATA GAA AGA GTC GCA GTG ACC GGA TCG CGA ATC GCT AAA GCA GAG CTA	T A A E E Q I E R V A V T G S R I A K A E L >		
9860	9880	9900	
ACT CAA CCA GCT CCA GTC GTC AGC CTT TCA GCC GAA CTG ACA AAA TTT GGT AAT CAA GAT TTA	T Q P A P V V S L S A E E L T K F G N Q D L >		
9920	9940	9960	
GGT AGC GTA CTA GCA GAA TTA CCT GCT ATT GGT GCA ACC AAC ACT ATT ATT GGT AAT AAC AAT AGC	G S V L A E L P A I G A T N T I I G N N N S >		
9980	10000	10020	10040
AAC TCA AGC GCA GGT GTT AGC TCA GCA GAC TTG CGT CGT CTA GGT GCT AAC AGA ACC TTA GTA TTA	N S S A G V S S A D L R R L G A N R T L V L >		
10060	10080	10100	
GTC AAC GGT AAG CGC TAC GTT GCC GGC CAA CCG GGC TCA GCT GAG GTA GAT TTG TCA ACT ATA CCA	V N G K R Y V A G Q P G S A E V D L S T I P >		
10120	10140	10160	
ACT AGC ATG ATC TCG CGA GTT GAG ATT GTA ACC GGC GGT GCT TCA GCA ATT TAT GGT TCG GAC GCT	T S M I S R V E I V T G G A S A I Y G S D A >		
10180	10200	10220	10240
GTA TCA CGT GTT ATC AAC GTT ATC CTT AAA GAA GAC TTT GAA GGC TTT GAG TTT AAC GCA CGT ACT	V S G V I N V I L K E D F E G F E F N A R T >		
10260	10280	10300	
AGC GGT TCT ACT GAA AGT GTA GGC ACT CAA GAG CAC TCT TTT GAC ATT TTG GGT GGT GCA AAC GTT	S G S T E S V G T Q E H S F D I L G G A N V >		
10320	10340	10360	
GCA GAT GGA CGT GGT AAT GTA ACC TTC TAC GCA GGT TAT GAA CGT ACA AAA GAA GTC ATG GCT ACC	A D G R G N V T F Y A G Y E R T K E V M A T >		
10380	10400	10420	
GAC ATT CGC CAA TTC GAT GCT TGG GGA ACA ATT AAA AAC GAA GCC GAT GGT GGT GAA GAT GAT GGT	D I R Q F D A W G T I K N E A D G G E D D G >		
10440	10460	10480	10500
ATT CCA GAC AGA CTA CGT GTA CCA CGA GGT TAT TCT GAA ATG ATT AAT GCT ACC GGT GTT ATC AAT	I P D R L R V P R V Y S E M I N A T G V I N >		

Fig. 4
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Fig. 4
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L R G T V G E A V R A P N I A E A F S P R S>
 11700 11720 11740
 CCT GGT TTT GGC CGC GTT TCA GAT CCA TGT GAT GCA GAT AAC ATT AAT GAC GAT CCG GAT CGC GTG
 P G F G R V S D P C D A D N I N D D P D R V>
 11760 11780 11800 11820
 TCA AAC TGT GCA GCA TTG GGG ATC CCT CCA GGA TTC CAA GCT AAT GAT AAC GTC AGT GTA GAT ACC
 S N C A A L G I P P G F Q A N D N V S V D T>
 11840 11860 11880
 TTA TCT GGT GGT AAC CCA GAT CTA AAA CCT GAA ACA TCA ACA TCC TTT ACA GGT GGT CTT GTT TGG
 L S G G N P D L K P E T S T S F T G G L V W>
 11900 11920 11940
 ACA CCA ACG TTT GCT GAC AAT CTA TCA TTC ACT GTC GAT TAT TAT GAT ATT CAA ATT GAG GAT GCT
 T P T F A D N L S F T V D Y Y D I Q I E D A>
 11960 11980 12000 12020
 ATT TTG TCA GTA GCC ACC CAG ACT GTG GCT GAT AAC TGT GTT GAC TCA ACT GGC GGA CCT GAC ACC
 I L S V A T Q T V A D N C V D S T G G P D T>
 12040 12060 12080
 GAC TTC TGT AGT CAA GTT GAT CGT AAT CCA ACG ACC TAT GAT ATT GAA CTT GTT CGC TCT GGT TAT
 D F C S Q V D R N P T T Y D I E L V R S G Y>
 12100 12120 12140
 CTA AAT GCC GCG GCA TTG AAT ACC AAA GGT ATT GAA TTT CAA GCT GCA TAC TCA TTA GAT CTA GAG
 L N A A A L N T K G I E F Q A A Y S L D L E>
 12160 12180 12200 12220
 TCT TTC AAC GCG CCT GGT GAA CTA CGC TTC AAC CTA TTG GGG AAC CAA TTA CTT GAA CTA GAA CGT
 S F N A P G E L R F N L L G N Q L L E L E R>
 12240 12260 12280
 CTT GAA TTC CAA AAT CGT CCT GAT GAG ATT AAT GAT GAA AAA GGC GAA GTA GGT GAT CCA GAG CTG
 L E F Q N R P D E I N D E K G E V G D P E L>
 12300 12320 12340
 CAG TTC CGC CTA GGC ATC GAT TAC CGT CTA GAT GAT CTA AGT GTT AGC TGG AAC ACG CGT TAT ATT
 Q F R L G I D Y R L D D L S V S W N T R Y I>
 12360 12380 12400
 GAT AGC GTA GTA ACT TAT GAT GTC TCT GAA AAT GGT GGC TCT CCT GAA GAT TTA TAT CCA GGC CAC
 D S V V T Y D V S E N G G S P E D L Y P G H>
 12420 12440 12460 12480
 ATA GGC TCA ATG ACA ACT CAT GAC TTG AGC GCT ACA TAC TAC ATC AAT GAG AAC TTC ATG ATT AAC
 I G S M T T H D L S A T Y Y I N E N F M I N>
 12500 12520 12540
 GGT GGT GTA CGT AAC CTA TTT GAC GCA CTT CCA CCT GGA TAC ACT AAC GAT GCG CTA TAT GAT CTA
 G G V R N L F D A L P P G Y T N D A L Y D L>
 12560 12580 12600 12620
 GTT CGT CGC CGT GCA TTC CTA GGT ATT AAG GTA ATG ATG TAATTAATTA TTACGCCTCT AACTAATAAA
 V G R R A F L G I K V M M>
 12640 12660 12680 12700
 AATGCAATCT CTTCTGAGAG ATTGCATTT TTTATGAAAT CCAATCTAA ACTGGTCTC CGAGCATCTT ACGCCCTAA
 12720 12740 12760 12780
 AACCCCCCCC CTCAATGTAA CGCCAAAGTT AATTGCTTAC ACGCACTTAC ACAAAACGAAC AATTTCATTA ACACGAGACA
 12800 12820 12840 12860
 CAGCTCACGC TTTTTATTTT ACCCTTGATT TTACTACATA AAATTGCGTT TTAGCCACA AGTGGTCTCC CAAGCTGGTC
 12880 12900 12920 12940
 GTATCTGTAA TTATTCAGTC CCAGGTGATT GTATTGACCC ATAAGCTCAG GTAGTCTGCT CTGCCATTAG CAAACAAATA
 12960 12980 13000 13020

Fig. 4
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TTGACAAAAT GGCATAAAA TGTGGCTTAG CGCTAAGTC ACCGTAAGTT TTATCGGCAT TAAGTCCAA CAGATTATTA
 13040 13060 13080
 ACGGAAACCC GCTAAACTG ATG GCA AAA ATA AAT ACT GAA CAC TTG GAT GAA GCT ACT ATT ACT TCG AAT
 M A K I N S E H L D E A T I T S N>
 13100 13120 13140
 AAG TGT ACG CAA ACA GAG ACT GAG GCT CGG CAT AGA AAT GCC ACT ACA ACA CCT GAG ATG CGC CGA
 K C T Q T E T E A R H R N A T T T P E M R R>
 13160 13180 13200 13220
 TTC ATA CAA GAG TCG GAT CTC AGT GTT AGC CAA CTG TCT AAA ATA TTA AAT ATC AGT GAA GCT ACC
 F I Q E S D L S V S Q L S K I L N I S E A T>
 13240 13260 13280
 GTA CGT AAG TGG CGC AAG CGT GAC TCT GTC GAA AAC TGT CCT AAT ACC CCG CAC CAT CTC AAT ACC
 V R K W R K R D S V E N C P N T P H H L N T>
 13300 13320 13340
 ACG CTA ACC CCT TTG CAA GAA TAT GTG GTT GTG GGC CTG CGT TAT CAA TTG AAA ATG CCA TTA GAC
 T L T P L Q E Y V V V G L R Y Q L K M P L D>
 13360 13380 13400 13420
 AGA TTG CTC AAA GCA ACC CAA GAG TTT ATC AAT CCA AAC GTG TCG CGC TCA GGT TTA GCA AGA TGT
 R L L K A T Q E F I N P N V S R S G L A R C>
 13440 13460 13480
 TTG AAG CGT TAT GGC CTT TCA CGG GTG AGT GAT ATC CAA ACC CCA CAC GTC CCA ATG CCC TAC TTT
 L K R Y G V S R V S D I Q S P H V P M R Y F>
 13500 13520 13540
 ATG CAA ATT CCA GTC ACT CAA GGC AGC GAT GTG CAA ACC TAC ACC CTG CAC TAT GAA ACG CTG GCA
 N Q I P V T Q G S D V Q T Y T L H Y E T L A>
 13560 13580 13600
 AAA ACC TTA GCC TTA CCT AGT ACC GAT GGT GAC AAT GTG GTG CAA GTG GTG TCT CTC ACC ATT CCA
 K T L A L P S T D G D N V V Q V V S L T I P>
 13620 13640 13660 13680
 CCA AAG TTA ACC GAA GAA GCA CCC AGT TCA ATT TTG CTC GGC ATT GAT CCT CAT AGC GAC TGG ATC
 P K L T E E A P S S I L L G I D P H S D W I>
 13700 13720 13740
 TAT CTC GAC ATA TAC CAA GAT GGC AAT ACA CAA GCC ACG AAT AGA TAT ATG GCT TAT GTG CTA AAA
 Y L D I Y Q D G N T Q A T N R Y M A Y V L K>
 13760 13780 13800
 CAC GGG CCA TTC CAT TTA CGA AAG TTA CTC GTG CGT AAC TAT CAC ACC TTT TTA CAG CGC TTT CCT
 H G P F H L R K L L V R N Y H T F L Q R F P>
 13820 13840 13860 13880
 GGA GCG ACG CAA AAT CGC CGC CCC TCT AAA GAT ATG CCT GAA ACA ATC AAC AAG ACG CCT GAA ACA
 G A T Q N R R P S K D M P E T I N K T P E T>
 13900 13920 13940
 CAG GCA CCC AGT GGA GAC TCA TA ATG AGC CAG ACC TCT AAA CCT ACA AAC TCA GCA ACT GAG CAA
 Q A P S G D S> M S Q T S K P T N S A T E Q>
 13960 13980 14000
 GCA CAA GAC TCA CAA GCT GAC TCT CGT TTA AAT AAA CGA CTA AAA GAT ATG CCA ATT GCT ATT GTT
 A Q D S Q A D S R L N K R L K D M P I A I V>
 14020 14040 14060
 GGC ATG GCG AGT ATT TTT GCA AAC TCT CGC TAT TTG AAT AAG TTT TGG GAC TTA ATC AGC GAA AAA
 G M A S I F A N S R Y L N K F W D L I S E K>
 14080 14100 14120 14140
 ATT GAT GCG ATT ACT GAA TTA CCA TCA ACT CAC TGG CAG CCT GAA GAA TAT TAC GAC GCA GAT AAA
 I D A I T E L P S T H W Q P E E Y Y D A D K>

O-f5

O-f6

Fig. 4
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14160 * 14180 * 14200 *
 ACC GCA GCA GAC AAA AGC TAC TGT AAA CGT GGT GGC TTT TTG CCA GAT GTA GAC TTC AAC CCA ATG
 T A A D K S Y C K R G G F L P D V D F N P M>
 14220 * 14240 * 14260 *
 GAG TTT GGC CTG CCG CCA AAC ATT TTG GAA CTG ACC GAT TCA TCG CAA CTA TTA TCA CTC ATC GTT
 E F G L P P N I L E L T D S S Q L L S L I V>
 14280 * 14300 * 14320 * 14340 *
 GCT AAA GAA GTG TTG GCT GAT GCT AAC TTA CCT GAG AAT TAC GAC CGC GAT AAA ATT GGT ATC ACC
 A K E V L A D A N L P E N Y D R D K I G I T>
 14360 * 14380 * 14400 *
 TTA GGT GTC GGC GGT GGT CAA AAA ATT AGC CAC AGC CTA ACA GCG CGT CTG CAA TAC CCA GTA TTG
 L G V G G G Q K I S H S L T A R L Q Y P V L>
 14420 * 14440 * 14460 *
 AAG AAA GTA TTC GCC AAT AGC GGC ATT AGT GAC ACC GAC AGC GAA ATG CTT ATC AAG AAA TTC CAA
 K K V F A N S G I S D T D S E M L I K K F Q>
 14480 * 14500 * 14520 * 14540 *
 GAC CAA TAT GTA CAC TGG GAA AAC TCG TTC CCA GGT TCA CTT GGT AAC GTT ATT GCG GGC CGT
 D Q Y V H W E E N S F P G S L G N V I A G R>
 14560 * 14580 * 14600 *
 ATC GCC AAC CGC TTC GAT TTT GGC GGC ATG AAC TGT GTG GTT GAT GCT GCC TGT GCT GGA TCA CTT
 I A N R F D F G G M N C V V D A A C A G S L>
 14620 * 14640 * 14660 *
 GCT GCT ATG CGT ATG GCG CTA ACA GAG CTA ACT GAA GGT CGC TCT GAA ATG ATG ATC ACC GGT GGT
 A A M R M A L T E L T E G R S E M M I T G G>
 14680 * 14700 * 14720 *
 GTG TGT ACT GAT AAC TCA CCC TCT ATG TAT ATG AGC TTT TCA AAA ACG CCC GCC TTT ACC ACT AAC
 V C T D N S P S M Y M S F S K T P A F T T N>
 14740 * 14760 * 14780 * 14800 *
 GAA ACC ATT CAG CCA TTT GAT ATC GAC TCA AAA GGC ATG ATG ATT GGT GAA GGT ATT GGC ATG GTG
 E T I Q P F D I D S K G M M I G E G I G M V>
 14820 * 14840 * 14860 *
 CCG CTA AAG CGT CTT GAA GAT GCA GAG CGC GAT GGC GAC CGC ATT TAC TCT GTA ATT AAA GGT GTG
 A L K R L E D A E R D G D R I Y S V I K G V>
 14880 * 14900 * 14920 *
 GGT GCA TCA TCT GAC GGT AAG TTT AAA TCA ATC TAT GGC CCT CGC CCA TCA GGC CAA GCT AAA CGA
 G A S S D G K F K S I Y A P R P S G Q A K A>
 14940 * 14960 * 14980 * 15000 *
 CTT AAC CGT GCC TAT GAT GAC GCA GGT TTT GGC CCG CAT ACC TTA GGT CTA ATT GAA GCT CAC GGA
 L N R A Y D D A G F A P H T L G L I E A H G>
 15020 * 15040 * 15060 *
 ACA GGT ACT GCA GCA GGT GAC GCG GCA GAG TTT GGC GGC CTT TGC TCA GTC TTT GCT GAA GGC AAC
 T G T A A G D A A E F A G L C S V F A E G N>
 15080 * 15100 * 15120 *
 GAT ACC AAG CAA CAC ATT GCG CTA CGT TCA GTT AAA TCA CAA ATT GGT CAT ACT AAA TCA ACT GCA
 D T K Q H I A L G S V K S Q I G H T K S T A>
 15140 * 15160 * 15180 * 15200 *
 GGT ACA GCA GGT TTA ATT AAA GCT GCT CTT GCT TTG CAT CAC AAG GTA CTG CCG CCG ACC ATT AAC
 G T A G L I K A A L A L H H K V L P P T I N>
 15220 * 15240 * 15260 *
 GTT AGT CAG CCA AGC CCT AAA CTT GAT ATC GAA AAC TCA CCG TTT TAT CTA AAC ACT GAG ACT CGT
 V S Q P S P K L D I E N S P F Y L N T E T R>
 15280 * 15300 * 15320 *
 CCA TGG TTA CCA CGT GTT GAT GGT ACG CCG CGC CGC GCG GGT ATT AGC TCA TTT GGT TTT GGT GGC
 P W L P R V D G T P R R A G I S S F G F G G>

Fig. 4
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15340 * 15360 * 15380 *
 ACT AAC TTC CAT TTT GTA CTA GAA GAG TAC AAC CAA GAA CAC AGC CGT ACT GAT AGC GAA AAA GCT
 T N F H F V L E E Y N Q E H S R T D S E K A>
 15400 * 15420 * 15440 * 15460 *
 AAG TAT CGT CAA CGC CAA GTG GCG CAA AGC TTC CTT GTT AGC GCA AGC GAT AAA GCA TCG CTA ATT
 K Y R Q R Q V A Q S F L V S A S D K A S L I>
 15480 * 15500 * 15520 *
 AAC GAG TTA AAC GTA CTA GCA GCA TCT GCA AGC CAA GCT GAG TTT ATC CTC AAA GAT GCA GCA GCA
 N E L N V L A A S A S Q A E F I L K D A A A A>
 15540 * 15560 * 15580 *
 AAC TAT GGC GTA CGT GAG CTT GAT AAA AAT GCA CCA CGG ATC GGT TTA GTT GCA AAC ACA GCT GAA
 N Y G V R E L D K N A P R I G L V A N T A E>
 15600 * 15620 * 15640 * 15660 *
 GAG TTA GCA GGC CTA ATT AAG CAA GCA CTT GCC AAA CTA GCA GCT AGC GAT GAT AAC GCA TGG CAG
 E L A G L I K Q A L A K L A A S D D N A W Q>
 15680 * 15700 * 15720 *
 CTA CCT GGT GGC ACT AGC TAC CGC GCC GCT GCA GTA GAA GGT AAA GTT GCC GCA CTG TTT GCT GGC
 L P G G T S Y R A A A V E G K V A A L F A G>
 15740 * 15760 * 15780 *
 CAA GGT TCA CAA TAT CTC AAT ATG GGC CGT GAC CTT ACT TGT TAT TAC CCA GAG ATG CGT CAG CAA
 Q G S Q Y L N M G R D L T C Y Y P E M R Q Q>
 15800 * 15820 * 15840 * 15860 *
 TTT GTA ACT GCA GAT AAA GTA TTT GCC GCA AAT GAT AAA ACG CCG TTA TCG CAA ACT CTG TAT CCA
 F V T A D K V F A A N D K T P L S Q T L Y P>
 15880 * 15900 * 15920 *
 AAG CCT GTA TTT AAT AAA GAT GAA TTA AAG GCT CAA GAA GCC ATT TTG ACC AAT ACC GCC AAT GCC
 K P V F N K D E L K A Q E A I L T N T A N A>
 15940 * 15960 * 15980 *
 CAA AGC GCA ATT GGT GCG ATT TCA ATG GGT CAA TAC GAT TTG TTT ACT GCG GCT GGC TTT AAT GCC
 Q S A I G A I S M G Q Y D L F T A A G F N A>
 16000 * 16020 * 16040 *
 GAC ATG GTT GCA GGC CAT AGC TTT GGT GAG CTA AGT GCA CTG TGT GCT GCA GGT GTT ATT TCA GCT
 D M V A G H S F G E L S A L C A A G V I S A>
 16060 * 16080 * 16100 * 16120 *
 GAT GAC TAC TAC AAG CTG GCT TTT GCT CGT GGT GAG GCT ATG GCA ACA AAA GCA CCG GCT AAA GAC
 D D Y Y K L A F A R G E A M A T K A P A K D>
 16140 * 16160 * 16180 *
 GGC GTT GAA GCA GAT GCA GGA GCA ATG TTT GCA ATC ATA ACC AAG AGT GCT GCA GAC CTT GAA ACC
 G V E A D A G A M F A I I T K S A A D L E T>
 16200 * 16220 * 16240 *
 GTT GAA GCC ACC ATC GCT AAA TTT GAT GGG GTG AAA GTC GCT AAC TAT AAC GCG CCA ACG CAA TCA
 V E A T I A K F D G V K V A N Y N A P T Q S>
 16260 * 16280 * 16300 * 16320 *
 GTA ATT GCA GGC CCA ACA GCA ACT ACC CCT GAT GCG GCT AAA GCG CTA ACT GAG CTT GGT TAC AAA
 V I A G P T A T T A D A A K A L T E L G Y K>
 16340 * 16360 * 16380 *
 GCG ATT AAC CTG CCA GTA TCA GGT GCA TTC CAC ACT GAA CTT GTT GGT CAC GCT CAA GCG CCA TTT
 A I N L P V S G A F H T E L V G H A Q A P F>
 16400 * 16420 * 16440 *
 GCT AAA GCG ATT GAC GCA GCC AAA TTT ACT AAA ACA AGC CGA GCA CTT TAC TCA AAT GCA ACT GGC
 A K A I D A A K F T K T S R A L Y S N A T G>
 16460 * 16480 * 16500 * 16520 *
 GGA CTT TAT GAA AGC ACT GCT GCA AAG ATT AAA GCC TCG TTT AAG AAA CAT ATG CTT CAA TCA GTG

Fig. 4
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Fig. 4
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TTC AGC CCC GAA ACA GCC CTG AGC GCA ACA AAA GTC CAA GCC ACT ATG CTT GAA GTG GTT GCT GAG
 F S A E T A L S A T K V Q A T M L E V V A E>
 * 17720 * 17740 * 17760 *
 * AAA ACC GGT TAC CCA ACT GAA ATG CTA GAG CTT GAA ATG GAT ATG GAA GCC GAT TTA GG GAT ATC GAT
 K T G Y P T E M L E M D M E A D L G I D>
 17780 * 17800 * 17820 * 17840 *
 * TCT ATC AAG CGT GAA ATT CTT GGT GCA ACA GAA GAT GAG CTA CCG GGT CTA CCT GAG CTT AGC
 S I K R V E I L G T V Q D E L P G L P E L S>
 * 17860 * 17880 * 17900 *
 * CCT GAA GAT CTA GCT GAG TGT CGA ACG CTA GGC GAA ATC GTT GAC TAT ATG GGC AGT AAA CTG CGG
 P E D L A E C R T L G E I V D Y M G S K L P>
 * 17920 * 17940 * 17960 *
 * GCT GAA GGC TCT ATG AAT TCT CAG CTG TCT ACA GGT TCC GCA GCT GCG ACT CCT GCA GCG AAT GGT
 A E G S M N S Q L S T G S A A A T P A A N G>
 17980 * 18000 * 18020 *
 * CTT TCT GCG GAG AAA GTT CAA GCG ACT ATG ATG TCT GTG GTT GCC GAA AAG ACT GGC TAC CCA ACT
 L S A E K V Q A T M M S V V A E K T G Y P T>
 18040 * 18060 * 18080 * 18100 *
 * GAA ATG CTA GAG CTT GAA ATG GAT ATG GAA GCC GAT TTA GGC ATA GAT TCT ATC AAG CGC GTT GAA
 E M L E M D M E A D L G I D S I K R V E>
 * 18120 * 18140 * 18160 *
 * ATT CTT GGC ACA GAA CAA GAT GAG CTA CCG GGT CTA CCT GAG CTT AGC CCT GAA GAT CTA GCT GAG
 I L G T V Q D E L P G L P E L S P E D L A E>
 18180 * 18200 * 18220 *
 * TGT CGT ACT CTA GGC GAA ATC GTT GAC TAT ATG AAC TCT AAA CTC GCT GAC GGC TCT AAG CTG CGG
 C R T L G E I V D Y M N S K L A D G S K L P>
 18240 * 18260 * 18280 * 18300 *
 * GCT GAA GGC TCT ATG AAT TCT CAG CTG TCT ACA AGT GCC GCA GCT GCG ACT CCT GCA GCG AAT GGT
 A E G S M N S Q L S T S A A A A T P A A N G>
 * 18320 * 18340 * 18360 *
 * CTC TCT GCG GAG AAA GTT CAA GCG ACT ATG ATG TCT GTG GTT GCC GAA AAG ACT GGC TAC CCA ACT
 L S A E K V Q A T M M S V V A E K T G Y P T>
 * 18380 * 18400 * 18420 *
 * GAA ATG CTA GAA CTT GAA ATG GAT ATG GAA GCT GAC CTT GGC ATC GAT TCA ATC AAG CGC GTT GAA
 E M L E M D M E A D L G I D S I K R V E>
 * 18440 * 18460 * 18480 * 18500 *
 * ATT CTT GGC ACA GAA CAA GAT GAG CTA CCG GGT TTA CCT GAG CTA AAT CCA GAA GAT TTG GCA GAG
 I L G T V Q D E L P G L P E L N P E D L A E>
 * 18520 * 18540 * 18560 *
 * TGT CGT ACT CTT GGC GAA ATC GTG ACT TAT ATG AAC TCT AAA CTC GCT GAC GGC TCT AAG CTG CGG
 C R T L G E I V T Y M N S K L A D G S K L P>
 * 18580 * 18600 * 18620 *
 * GCT GAA GGC TCT ATG CAC TAT CAG CTG TCT ACA AGT ACC GCT GCT GCG ACT CCT GCA GCG AAT GGT
 A E G S M H Y Q L S T S T A A A A T P V A N G>
 * 18640 * 18660 * 18680 *
 * CTC TCT GCA GAA AAA GTT CAA GCG ACC ATG ATG TCT GTG GTT GCA GAT AAA ACT GGC TAC CCA ACT
 L S A E K V Q A T M M S V V A D K T G Y P T>
 18700 * 18720 * 18740 * 18760 *
 * GAA ATG CTT GAA CTT GAA ATG GAT ATG GAA GCC GAT TTA GGT ATC GAT TCT ATC AAG CGC GTT GAA
 E M L E M D M E A D L G I D S I K R V E>
 * 18780 * 18800 * 18820 *
 * ATT CTT GGC ACA GAA CAA GAT GAG CTA CCG GGT TTA CCT GAG CTA AAT CCA GAA GAT CTA GCA GAG
 I L G T V Q D E L P G L P E L N P E D L A E>
 * 18840 * 18860 * 18880 *

Fig. 4
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MISSING AT THE TIME OF PUBLICATION

Fig. 4
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21220 * 21240 * 21260 *
 GGT ACG TCA ATG CAA GGT GGC AGC GAC ACT AAA GCA ACT GAG ACT GCT TCT GTA AAA AAG CTT AAT
 G T S M Q G G S D T K A T E T A S V K K L N>
 21280 * 21300 * 21320 *
 GCG GGT GAG GTG CTA AGT GCA TCG CAT CCG CGT GCT GGT GCA CAA AAA ACA CCA CTA CAA GCT GTC
 A G E V L S A S H P R A G A Q K T P L Q A V>
 21340 * 21360 * 21380 * 21400 *
 ACT GCA ACG CGT CTG TTA ACC CCA AGT GCC ATG GTC TTC ATT GAA GAT CAC CGC ATT GGC GGT AAC
 T A T R L L T P S A M V F I E D H R I G G N>
 21420 * 21440 * 21460 *
 AGT GTG TTG CCA ACG GTA TGC GCC ATC GAC TGG ATG CGT GAA GCG GCA ACC GAC ATG CTT GGC GCT
 S V L P T V C A I D W M R E A A S D M L G A>
 21480 * 21500 * 21520 *
 CAA GTT AAG GTA CTT GAT TAC AAG CTA TTA AAA GGC ATT GTA TTT GAG ACT GAT GAG CCG CAA GAG
 Q V K V L D Y K L L K G I V F E T D E P Q E>
 21540 * 21560 * 21580 * 21600 *
 TTA ACA CTT GAG CTA ACG CCA GAC GAT TCA GAC GAA GCT ACG CTA CAA GCA TTA ATC AGC TGT AAT
 L T L E L T P D D S D E A T L Q A L I S C N>
 21620 * 21640 * 21660 *
 GGG CGT CCG CAA TAC AAG GCG ACG CTT ATC AGT GAT AAT GCC GAT ATT AAG CAA CTT AAC AAG CAG
 G R P Q Y K A T L I S D N A D I K Q L N K Q>
 21680 * 21700 * 21720 *
 TTT GAT TTA AGC GCT AAG GCG ATT ACC ACA GCA AAA GAG CTT TAT AGC AAC GGC ACC TTG TTC CAC
 F D L S A K A I T T A K E L Y S N G T L F H>
 21740 * 21760 * 21780 * 21800 *
 GGT CCG CGT CTA CAA GGG ATC CAA TCT GTA GTG CAG TTC GAT GAT CAA GGC TTA ATT GCT AAA GTC
 G P R L Q G I Q S V V Q F D D Q G L I A K V>
 21820 * 21840 * 21860 *
 GCT CTG CCT AAG GTT GAA CTT AGC CAT TGT GGT GAG TTC TTG CCG CAA ACC CAC ATG GGT GGC AGT
 A L P K V E L S D C G E F L P Q T H M G G S>
 21880 * 21900 * 21920 *
 CAA CCT TTT GCT GAG GAC TTG CTA TTA CAA GCT ATG CTG GTT TGG GCT CGC CTT AAA ACT GGC TCG
 Q P F A E D L L Q A M L V W A R L K T G S>
 21940 * 21960 * 21980 *
 GCA AGT TTG CCA TCA AGC ATT GGT GAG TTT ACC TCA TAC CAA CCA ATG GCC TTT GGT GAA ACT GGT
 A S L P S S I G E F T S Y Q P M A F G E T G>
 22000 * 22020 * 22040 * 22060 *
 ACC ATA GAG CTT GAA GTG ATT AAG CAC AAC AAA CGC TCA CTT GAA GCG AAT GTT GCG CTA TAT CGT
 T I E L E V I K H N K R S L E A N V A L Y R>
 22080 * 22100 * 22120 *
 GAC AAC GGC GAG TTA AGT CCC ATG TTT AAG TCA GCT AAA ATC ACC ATT AGC AAA ACC TTA AAT TCA
 D N G E L S A M F K S A K I T I S K S L N S>
 22140 * 22160 * 22180 * 22200 *
 GCA TTT TTA CCT GCT GTC TTA GCA AAC GAC AGT GAG GCG AAT TAGTGGAA ACAAAACGCCT AAAGCTAGTG
 A F L P A V L A N D S E A N>
 22220 * 22240 * 22260 *
 CG ATG CCG CTG CGC ATC GCA CTT ATC TTA CTG CCA ACA CCG CAG TTT GAA GTT AAC TCT GTC GAC
 M P L R I A L I L P T P Q F E V N S V D>
 22280 * 22300 * 22320 *
 CAG TCA GTA TTA GCC AGC TAT CAA ACA CTG CAG CCT GAG CTA AAT GCC CTG CTT AAT AGT GCG CCG
 Q S V L A S Y Q T L Q P E L N A L L N S A P>
 22340 * 22360 * 22380 *
 ACA CCT GAA ATG CTC AGC ATC ACT ATC TCA GAT GAT AGC GAT GCA AAC AGC TTT GAG TCG CAG CTA

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Fig. 4
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T	P	E	M	L	S	I	T	I	S	D	D	S	D	A	N	S	F	E	S	Q	L>
22400	*	*	*	22420	*	*	*	*	22440	*	*	*	*	*	22460	*	*	*	*	*	*
AAT GCT GCG ACC AAC GCA ATT AAC AAT GGC TAT ATC GTC AAG CTT GCT ACG GCA ACT CAC GCT TTG N A A T N A I N N G Y I V K L A T A T H A L>																					
22480	*	*	*	22500	*	*	*	*	22520	*	*	*	*	*	*	*	*	*	*	*	
TTA ATG CTG CCT GCA TTA AAA GCG GCG CAA ATG CGG ATC CAT CCT CAT GCG CAG CTT GCC GCT ATG L M L P A L K A A Q M R I H P H A Q L A A M>																					
22540	*	*	*	22560	*	*	*	*	22580	*	*	*	*	*	*	*	*	*	*	*	
CAG CAA GCT AAA TCG ACG CCA ATG AGT CAA GTA TCT GGT GAG CTA AAG CTT GGC GCT AAT GCG CTA Q Q A K S T P M S Q V S G E L K L G A N A L>																					
22600	*	*	*	22620	*	*	*	*	22640	*	*	*	*	*	22660	*	*	*	*	*	
AGC CTA GCT CAG ACT AAT GCG CTG TCT CAT GCT TTA AGC CAA GCC AAG CGT AAC TTA ACT GAT GTC S L A Q T N A L S H A L S Q A K R N L T D V>																					
22680	*	*	*	22700	*	*	*	*	22720	*	*	*	*	*	*	*	*	*	*	*	
AGC GTG AAT GAG TGT TTT GAG AAC CTC AAA AGT GAA CAG CAG TTC ACA GAG GTT TAT TCG CTT ATT S V N E C F E N L K S E Q Q F T E V Y S L I>																					
22740	*	*	*	22760	*	*	*	*	22780	*	*	*	*	*	*	*	*	*	*	*	
CAG CAA CTT GCT AGC CGC ACC CAT GTG AGA AAA GAG GTT AAT CAA GGT GTG GAA CTT GCC CCT AAA Q Q L A S R T H V R K E V N Q G V E L G P K>																					
22800	*	*	*	22820	*	*	*	*	22840	*	*	*	*	*	*	*	*	*	*	*	
CAA GCC AAA AGC CAC TAT TGG TTT AGC GAA TTT CAC CAA AAC CGT GTT GCT GCC ATC AAC TTT ATT Q A K S H Y W F S E F H Q N R V A A I N F I>																					
22860	*	*	*	22880	*	*	*	*	22900	*	*	*	*	*	22920	*	*	*	*	*	
AAT GGC CAA CAA GCA ACC AGC TAT GTG CTT ACT CAA GGT TCA GGA TTG TTA GCT GCG AAA TCA ATG N G Q Q A T S Y V L T Q G S G L L A A K S M>																					
22940	*	*	*	22960	*	*	*	*	22980	*	*	*	*	*	*	*	*	*	*	*	
CTA AAC CAG CAA AGA TTA ATG TTT ATC TTG CCG GGT AAC AGT CAG CAA CAA ATA ACC GCA TCA ATA L N Q Q R L M F I L P G N S Q Q Q I T A S I>																					
23000	*	*	*	23020	*	*	*	*	23040	*	*	*	*	*	*	*	*	*	*	*	
ACT CAG TTA ATG CAG CAA TTA GAG CGT TTG CAG GTA ACT GAG GTT AAT GAG CTT TCT CTA GAA TGC T Q L M Q Q L E R L Q V T E V N E L S L E C>																					
23060	*	*	*	23080	*	*	*	*	23100	*	*	*	*	*	23120	*	*	*	*	*	
CAA CTA GAG CTG CTC AGC ATA ATG TAT GAC AAC TTA GTC AAC GCA GAC AAA CTC ACT ACT CGC GAT Q L E L L S I M Y D N L V N A D K L T T R D>																					
23140	*	*	*	23160	*	*	*	*	23180	*	*	*	*	*	*	*	*	*	*	*	
AGT AAG CCC GCT TAT CAG GCT GTG ATT CAA GCA AGC TCT GTT AGC GCT GCA AAG CAA GAG TTA AGC S K P A Y Q A V I Q A S S V S A A K Q E L S>																					
23200	*	*	*	23220	*	*	*	*	23240	*	*	*	*	*	*	*	*	*	*	*	
GCG CTT AAC GAT GCA CTC ACA GCG CTG TTT GCT GAG CAA ACA AAC GCC ACA TCA ACG AAT AAA GGC A L N D A L T A L F A E Q T N A T S T N K G>																					
23260	*	*	*	23280	*	*	*	*	23300	*	*	*	*	*	23320	*	*	*	*	*	
TTA ATC CAA TAC AAA ACA CCG GCG AGT TAC TTA ACC CTA ACA CCG CTT GGC AGC AAC AAT GAC L I Q Y K T P A G S Y L T L T P L G S N N D>																					
23340	*	*	*	23360	*	*	*	*	23380	*	*	*	*	*	*	*	*	*	*	*	
AAC GCC CAA GCG GGT CTT GCT TTT GTC TAT CCG GGT GTG GGA ACG GTT TAC GCC GAT ATG CTT AAT N A Q A G L A F V Y P G V G T V Y A D M L N>																					
23400	*	*	*	23420	*	*	*	*	23440	*	*	*	*	*	*	*	*	*	*	*	
GAG CTG CAT CAG TAC TTC CCT GCG CTT TAC GCC AAA CTT GAG CGT GAA GGC GAT TTA AAG GCG ATG E L H Q Y F P A L Y A K L E R E G D L K A M>																					
23460	*	*	*	23480	*	*	*	*	23500	*	*	*	*	*	*	*	*	*	*	*	
CTA CAA GCA GAA GAT ATC TAT CAT CTT GAC CCT AAA CAT GCT GCC CAA ATG AGC TTA GCT GAC TTA L Q A E D I Y H L D P K H A A Q M S L G D L>																					
23520	*	*	*	23540	*	*	*	*	23560	*	*	*	*	*	23580	*	*	*	*	*	

Fig. 4
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GCC ATT GCT GGC GTG CGG AGC AGC TAC CTG TTA ACT CAG CTG CTC ACC GAT GAG TTT AAT ATT AAG
 A I A G V G S S Y L L T Q L L T D E F N I K>
 * 23600 * 23620 * 23640 *
 CCT AAT TTT GCA TTA GGT TAC TCA ATG GGT GAA GCA TCA ATG TGG GCA AGC TTA GGC GTA TGG CAA
 P N F A L G Y S M G E A S M W A S L G V W Q>
 * 23660 * 23680 * 23700 *
 AAC CCG CAT GCG CTG ATC AGC AAA ACC CAA ACC GAC CCG CTA TTT ACT TCT GCT ATT TCC GGC AAA
 N P H A L I S K T Q D P L F T S A I S G K>
 23720 * 23740 * 23760 * 23780 *
 TTG ACC GCG GTT AGA CAA GCT TGG CAG CTT GAT GAT ACC GCA GCG GAA ATC CAG TGG AAT AGC TTT
 L T A V R Q A W Q L D D T A A E I Q W N S F>
 * 23800 * 23820 * 23840 *
 GTG GTT AGA AGT GAA GCA GCG CCG ATT GAA GCC TTG CTA AAA GAT TAC CCA CAC GCT TAC CTC GCG
 V V R S E A A P I E A L L K D Y P H A Y L A>
 * 23860 * 23880 * 23900 *
 ATT ATT CAA GGG GAT ACC TGC GTA ATC GCT GGC TGT GAA ATC CAA TGT AAA GCG CTA CTT GCA GCA
 I I Q G D T C V I A G C E I Q C K A L L A A>
 23920 * 23940 * 23960 * 23980 *
 CTG GGT AAA CGC GGT ATT GCA GCT AAT CGT GTA ACG GCG ATG CAT ACG CAG CCT GCG ATG CAA GAG
 L G K R G I A A N R V T A M H T Q P A M Q E>
 * 24000 * 24020 * 24040 *
 CAT CAA AAT GTG ATG GAT TTT TAT CTG CAA CCG TTA AAA GCA GAG CTT CCT AGT GAA ATA AGC TTT
 H Q N V M D F Y L Q P L K A E L P S E I S F>
 * 24060 * 24080 * 24100 *
 ATC AGC GCC GCT GAT TTA ACT GCC AAG CAA ACY GTG AGT GAG CAA GCA CTT AGC AGC CAA GTC GTT
 I S A A D L T A K Q T V S E Q A L S S Q V V>
 * 24120 * 24140 * 24160 *
 GCT CAG TCT ATT GCC GAC ACC TTC TGC CAA ACC TTG GAC TTT ACC GCG CTA GTA CAT CAC GCC CAA
 A Q S I A D T F C Q T L D F T A L V H H A Q>
 24180 * 24200 * 24220 * 24240 *
 CAT CAA GGC GCT AAG CTG TTT GTT GAA ATT GCC GCG GAT AGA CAA AAC TGC ACC TTG ATA GAC AAG
 H Q G A K L F V E I G A D R Q N C T L I D K>
 * 24260 * 24280 * 24300 *
 ATT GTT AAA CAA GAT GGT GCC AGC ACT GTA CAA CAT CAA CCT TGT TGC ACA GTG CCT ATG AAC GCA
 I V K Q D G A S S V Q H Q P C C T V P M N A>
 * 24320 * 24340 * 24360 *
 AAA GGT AGC CAA GAT ATT ACC AGC GTG ATT AAA GCG CTT GGC CAA TTA ATT AGC CAT CAG GTG CCA
 K G S Q D I T S V I K A L G Q L I S H Q V P>
 24380 * 24400 * 24420 * 24440 *
 TTA TCG GTG CAA CCA TTT ATT GAT GGA CTC AAG CGC GAG CTA ACA CTT TGC CAA TTG ACC AGC CAA
 L S V Q P F I D G L K R E L T L C Q L T S Q>
 * 24460 * 24480 * 24500 *
 CAG CTG GCA GCA CAT GCA AAT GTT GAC AGC AAG TTT GAG TCT AAC CAA GAC CAT TTA CTT CAA GGG
 Q L A A H A N V D S K F E S N Q D H L L Q G>
 * 24520 * 24540 * 24560 *
 GAA GTC TA ATG TCA TTA CCA GAC AAT GCT TCT AAC CAC CTT TCT GCC AAC CAG AAA GGC GCA TCT
 E V>
 * 24580 * 24600 * 24620 * 24640 *
 CAG GCA AGT AAA ACC AGT AAG CAA AGC AAA ATC GCC ATT GTC GGT TTA GCC ACT CTG TAT CCA GAC
 Q A S K T S K Q S K I A I V G L A T L Y P D>
 * 24660 * 24680 * 24700 *
 GCT AAA ACC CCG CAA GAA TTT TGG CAG AAT TTG CTG GAT AAA CGC GAC TCT CGC AGC ACC TTA ACT
 A K T P Q E F W Q N L L D K R D S R S T L T>

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Fig. 4
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KAS Tum

25900 * 25920 * 25940 * 25960 *

CCA TCG AAT AAT CAT TTT GGT GTA AGA ACC CGT CAC GCA GGC GTA TCG GTA TTT GGC TTT GGT GGC
P S N N H F G V R T R H A G V S V F G F G G>

25980 * 26000 * 26020 *

TGT AAC GCC CAT CTG TTG CTT GAG TCA TAC AAC GGC AAA GGA ACA GTA AAG GCA GAA GCC ACT CAA
C N A H L L E S Y N G K G T V K A E A T Q>

26040 * 26060 * 26080 *

GTA CCG CGT CAA GCT GAG CCG CTA AAA GTG GTT GGC CTT GCC TCG CAC TTT GGG CCT CTT AGC AGC
V P R Q A E P L K V V G L A S H F G P L S S>

26100 * 26120 * 26140 *

AT T AAT GCA CTC AAC AAT GCT GTG ACC CAA GAT GGG AAT GGC TTT ATC GAA CTG CCG AAA AAG CGC
I N A L N N A V T Q D G N G F I E L P K K R>

26160 * 26180 * 26200 * 26220 *

TGG AAA GGC CTT GAA AAG CAC AGT GAA CTG TTA GCT GAA TTT GGC TTA GCA TCT GCG CCA AAA GGT
W K G L E K H S E L L A E F G L A S A P K G>

26240 * 26260 * 26280 *

GCT TAT GTT GAT AAC TTC GAG CTG GAC TTT TTA CGC TTT AAA CTG CCG CCA AAC GAA GAT GAC CGT
A Y V D N F E L D F L R F K L P P N E D D R>

26300 * 26320 * 26340 *

TTG ATC TCA CAG CAG CTA ATG CTA ATG CGA GTC ACA GAC GAA GCC ATT CGT GAT GCC AAG CTT GAG
L I S Q Q L M L M R V T D E A I R D A K L E>

26360 * 26380 * 26400 * 26420 *

CCG GGG CAA AAA GTA GCT GTA TTA GTG GCA ATG GAA ACT GAG CTT GAA CTG CAT CAG TTC CGC GGC
P G Q K V A V L V A M E T E L E L H Q F R G>

26440 * 26460 * 26480 *

CGG GTT AAC TTG CAT ACT CAA TTA GCG CAA AGT CTT GCC GCC ATG GGC GTG AGT TTA TCA ACG GAT
R V N L H T Q L A Q S L A A M G V S L S T D>

26500 * 26520 * 26540 *

GAA TAC CAA GCG CTT GAA GCC ATC GCC ATG GAC AGC GTG CTT GAT GCT GCC AAG CTC AAT CAG TAC
E Y Q A L E A I A M D S V L D A A K L N Q Y>

26560 * 26580 * 26600 * 26620 *

ACC AGC TTT ATT GGT AAT ATT ATG GCG TCA CGC GTG GCG TCA CTA TGG GAC TTT AAT GGC CCA GCC
T S F I G N I M A S R V A S L W D F N G P A>

26640 * 26660 * 26680 *

TTC ACT ATT TCA GCA GCA GAG CAA TCT GTG AGC CGC TGT ATC GAT GTG GCG CAA AAC CTC ATC ATG
F T I S A A E Q S V S R C I D V A Q N L I M>

26700 * 26720 * 26740 *

GAG GAT AAC CTA GAT GCG GTG GTG ATT GCA GCG GTC GAT CTC TCT GGT AGC TTT GAG CAA GTC ATT
E D N L D A V V I A A V D L S G S F E Q V I>

26760 * 26780 * 26800 *

CTT AAA AAT GCC ATT GCA CCT GTA GCC ATT GAG CCA AAC CTC GAA GCA AGC CTT AAT CCA ACA TCA
L K N A I A P V A I E P N L E A S L N P T S>

26820 * 26840 * 26860 * 26880 *

GCA AGC TGG AAT GTC GGT GAA GGT GCT GGC GCG GTC GTG CTT GTT AAA AAT GAA GCT ACA TCG GGC
A S W N V G E G A G A V V L V K N E A T S G>

26900 * 26920 * 26940 *

TGC TCA TAC GGC CAA ATT GAT GCA CTT GGC TTT GCT AAA ACT GCC GAA ACA GCG TTG GCT ACC GAC
C S Y G Q I D A L G F A K T A E T A L A T D>

26960 * 26980 * 27000 *

AAG CTA CTG AGC CAA ACT GCC ACA GAC TTT AAT AAG GTT AAA GTG ATT GAA ACT ATG GCA GCG CCT
K L L S Q T A T D F N K V K V I E T M A A P>

27020 * 27040 * 27060 * 27080 *

GCT AGC CAA ATT CAA TTA GCG CCA ATA GTT AGC TCT CAA GTG ACT CAC ACT GCT GCA GAG CAG CGT

Fig. 4
21/30

Fig. 4
22/30

ATC CCT GTT GAT GCG CCG TAC TTA GTA GAC GGA CAA ATC CCT TGG GCG GTA GCA GTA GAA TCA GGC
 I P V D A P Y L V D G Q I P W A V A V E S G>
 28280 28300 28320
 CAA TGT GAC TTG ATG CTT ATT AGC TAT CTC GGT ATC GAC TTT GAG AAC AAA GGC GAG CGG GTT TAT
 Q C D L M L I S Y L G I D F E N K G E R V Y>
 28340 28360 28380 28400
 CGA CTA CTC GAT TGT ACC CTC ACC TTC CTA GGC GAC TTG CCA CGT GGC GGA GAT ACC CTA CGT TAC
 R L L D C T L T F L G D L P R G G D T L R Y>
 28420 28440 28460
 GAC ATT AAG ATC AAT AAC TAT GCT CGC AAC GGC GAC ACC CTG CTG TTC TTC TCG TAT GAG TGT
 D I K I N N Y A R N G D T L L F F F S Y E C>
 28480 28500 28520
 TTT GTT GGC GAC AAG ATG ATC CTC AAG ATG GAT GGC GGC TGC GCT GGC TTC TTC ACT GAT GAA GAG
 F V G D K M I L K M D G G C A G F F T D E E>
 28540 28560 28580 28600
 CTT GCC GAC GGT AAA GGC GTG ATT CGC ACA GAA GAA GAG ATT AAA GCT CGC AGC CTA GTG CAA AAG
 L A D G K G V I R T E E E I K A R S L V Q K>
 28620 28640 28660
 CAA CGC TTT AAT CCG TTA CTA GAT TGT CCT AAA ACC CAA TTT AGT TAT GGT GAT ATT CAT AAG CTA
 Q R F N P L L D C P K T Q F S Y G D I H K L>
 28680 28700 28720
 TTA ACT GCT GAT ATT GAG GGT TGT TTT GGC CCA AGC CAC AGT GGC GTC CAC CAG CCG TCA CTT TGT
 L T A D I E G C F G P S H S G V H Q P S L C>
 28740 28760 28780
 TTC GCA TCT GAA AAA TTC TTG ATG ATT GAA CAA GTC AGC AAG GTT GAT CGC ACT GGC GGT ACT TGG
 F A S E K F L M I E Q V S K V D R T G G T W>
 28800 28820 28840 28860
 GGA CTT GGC TTA ATT GAG GGT CAT AAG CAG CTT GAA GCA GAC CAC TGG TAC TTC CCA TGT CAT TTC
 G L G L I E G H K Q L E A D H W Y F P C H F>
 28880 28900 28920
 AAG GGC GAC CAA GTG ATG GCT GGC TCG CTA ATG GCT GAA GGT TGT GGC CAG TTA TTG CAG TTC TAT
 K G D Q V M A G S L M A E G C G Q L L Q F Y>
 28940 28960 28980
 ATG CTG CAC CTT GGT ATG CAT ACC CAA ACT AAA AAT GGT CGT TTC CAA CCT CTT GAA AAC GGC TCA
 M L H L G M H T Q T K N G R F Q P L E N A S>
 29000 29020 29040 29060
 CAG CAA GTA CGC TGT CGC GGT CAA GTG CTG CCA CAA TCA GGC GTG CTA ACT TAC CGT ATG GAA GTG
 Q Q V R C R G Q V L P Q S G V L T Y R M E V>
 29080 29100 29120
 ACT GAA ATC GGT TTC AGT CCA CGC CCA TAT GCT AAA GCT AAC ATC GAT ATC TTG CTT AAT GGC AAA
 T E I G F S P R P Y A K A N I D I L L N G K>
 29140 29160 29180
 GCG GTA GTG GAT TTC CAA AAC CTA GGG GTG ATG ATA AAA GAG GAA GAT GAG TGT ACT CGT TAT CCA
 A V V D F Q N L G V M I K E E D E C T R Y P>
 29200 29220 29240 29260
 CTT TTG ACT GAA TCA ACA ACG GCT AGC ACT GCA CAA GTA AAC GCT CAA ACA AGT CCG AAA AAG GTA
 L L T E S T T A S T A Q V N A Q T S A K K V>
 29280 29300 29320
 TAC AAG CCA GCA TCA GTC ATT GCG CCA TTA ATG GCA CAA ATT CCT GAT CTG ACT AAA GAG CCA AAC
 Y K P A S V N A P L M A Q I P D L T K E P N>
 29340 29360 29380
 AAG GGC GTT ATT CCG ATT TCC CAT GTT GAA GCA CCA ATT ACG CCA GAC TAC CCG AAC CGT GTA CCT
 K G V I P I S H V E A P I T P D Y P N R V P>
 29400 29420 29440

Fig. 4
23/30

GAT ACA GTG CCA TTC ACG CCG TAT CAC ATG TTT GAG TTT GCT ACA GGC AAT ATC GAA AAC TGT TTC
 D T V P F T P Y H M F E F A T G N I S N C F>
 29460 9480 29500 29520
 GGG CCA GAG TTC TCA ATC TAT CGC GGC ATG ATC CCA CCA CGT ACA CCA TCC GGT GAC TTA CAA GTG
 G P E F S I Y R G M I P P R T P C G D L Q V>
 29540 29560 29580
 ACC ACA CGT GTG ATT GAA GTT AAC GGT AAG CGT GGC GAC TTT AAA AAG CCA TCA TCG TGT ATC GCT
 T T R V I E V N G K R G D F K K P S S C I A>
 29600 29620 29640
 GAA TAT GAA GTG CCT GCA GAT GCG TGG TAT TTC GAT AAA AAC AGC CAC GGC GCA GTG ATG CCA TAT
 E Y E V P A D A W Y F D K N S H G A V M P Y>
 29660 29680 29700 29720
 TCA ATT TTA ATG GAG ATC TCA CTG CAA CCT AAC GGC TTT ATC TCA GGT TAC ATG GGC ACA ACC CTA
 S I L M E I S L Q P N G F I S G Y M G T T L>
 29740 29760 29780
 GGC TTC CCT GGC CTT GAG CTG TTC CGT AAC TTA GAC GGT AGC GGT GAG TTA CTA CGT GAA GTA
 G F P G L E L F F R N L D G S G E L L R E V>
 29800 29820 29840
 GAT TTA CGT GGT AAA ACC ATC CGT AAC GAC TCA CGT TTA TTA TCA ACA GTG ATG GCC ACT AAC
 D L R G K T I R N D S R L L S T V M A G T N>
 29860 29880 29900 29920
 ATC ATC CAA AGC TTT AGC TTC GAG CTA AGC ACT GAC GGT GAG CCT TTC TAT CGC GCC ACT GCG GTA
 I I Q S F S F E L S T D G E P F Y R G T A V>
 29940 29960 29980
 TTT GGC TAT TTT AAA GGT GAC GCA CTT AAA GAT CAG CTA GGC CTA GAT AAC GGT AAA GTC ACT CAG
 F G Y F K G D A L K D Q L G L D N G K V T Q>
 30000 30020 30040
 CCA TGG CAT GTA GCT AAC GGC GTT GCT GCA AGC ACT AAG GTG AAC CTG CTT GAT AAG AGC TGC CGT
 P W H V A N G V A A S T K V N L L D K S C R>
 30060 30080 30100
 CAC TTT AAT GCG CCA GCT AAC CAG CCA CAC TAT CGT CTA GCC GGT CAG CTG AAC TTT ATC GAC
 H F N A P A N Q P H Y R L A G G Q L N F I D>
 30120 30140 30160 30180
 AGT GTT GAA ATT GTT GAT AAT GGC GGC ACC GAA GGT TTA GGT TAC TTG TAT GCC GAG CGC ACC ATT
 S V E I V D N G G T E G L G Y L Y A E R T I>
 30200 30220 30240
 GAC CCA AGT GAT TGG TTC CAG TTC CAC CAC CAA GAT CCG GTT ATG CCA GGC TCC TTA GGT
 D P S D W F F Q F H F H Q D P V M P G S L G>
 30260 30280 30300
 GTT GAA GCA ATT ATT GAA ACC ATG CAA GCT TAC GCT ATT AGT AAA GAC TTG GGC GCA GAT TTC AAA
 V E A I I E T M Q A Y A I S K D L G A D F K>
 30320 30340 30360 30380
 AAT CCT AAG TTT CGT CAG ATT TTA TCG AAC ATC AAG TGG AAG TAT CGC GGT CAA ATC AAT CGG CTG
 N P K F G Q I L S N I K W K Y R G Q I N P L>
 30400 30420 30440
 AAC AAG CAG ATG TCT ATG GAT GTC AGC ATT ACT TCA ATC AAA GAT GAA GAC GGT AAG AAA GTC ATC
 N K Q M S M D V S I T S I K D E D G K K V I>
 30460 30480 30500
 ACA GGT AAT GCC AGC TTG AGT AAA GAT GGT CTG CGC ATA TAC GAG GTC TTC GAT ATA GCT ATC AGC
 T G N A S L S K D G L R I Y E V F D I A I S>
 30520 30540 30560 30580
 ATC GAA GAA TCT GTA T AAATCGGAGT GACTGTCTGG CTATTTACT CAATTTCTGT GTCAAAAGTG CTCACCTATA
 I E E S V>

Fig. 4
24/30

0.89

Fig. 4
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Fig. 4
26/30

33140 * 33160 * 33180 * 33200 *
 AGT AAG CCG TTA GAC TCC CCT GAT GTG CCT TCT ACC CAT GGG GTT ATC GCC ACA CGA TAC GGT
 S K P L D S P D D V P S T H G V I A T R Y G>
 33220 * 33240 * 33260 *
 CCA GCA ATT TAT AGC TCT ACC AGC ATT TTA AAA TCT GAT CGT AGC GGC TCC CAA CTT GGT TAT TTA
 P A I Y S S T S I L K S D R S G S Q L G Y L>
 33280 * 33300 * 33320 *
 GTC TTC ATT AGG TTA ATT GAT GAA TGG TTC ATC GCT GAG CTA TCG CAA TAC ACT GCC GCA GGT GTT
 V F I R L I D E W F I A E L S Q Y T A A G V>
 33340 * 33360 * 33380 * 33400 *
 GAA ATC GCT ATG GCT GAT GCC GCA GAC GCA CAA TTA GCG AGA TTA GGC GCA AAC ACT AAG CTT AAT
 E I A M A D A A D A Q L A R L G A N T K L N>
 33420 * 33440 * 33460 *
 AAA GTA ACC GCT ACA TCC GAA CGG TTA ATA ACT AAT GTC GAT GGT AAG CCT CTG TTG AAG TTA GTG
 K V T A T S E R L I T N V D G K P L L K L V>
 33480 * 33500 * 33520 *
 CTT TAC CAT ACC AAT AAC CAA CGG CCG CCG ATG CTA GAT TAC AGT ATA ATA ATT CTA TTA GTT GAG
 L Y H T N N Q P P P M L D Y S I I L L V E>
 33540 * 33560 * 33580 *
 ATG TCA TTT TTA CTG ATC CTC GCT TAT TTC CTT TAC TCC TAC TTC TTA GTC AGG CCA GTT AGA AAG
 M S F L L I L A Y F L Y S Y F L V R P V R K>
 33600 * 33620 * 33640 * 33660 *
 CTG GCT TCA GAT ATT AAA AAA ATG GAT AAA AGT CGT CAA ATT AAA AAG CTA AGG TAT CAC TAC CCT
 L A S D I K K M D K S R E I K K L R Y H Y P>
 33680 * 33700 * 33720 *
 ATT ACT GAG CTA GTC AAA GTT GCG ACT CAC TTC AAC GCC CTA ATG GGG ACG ATT CAG GAA CAA ACT
 I T E L V K V A T H F N A L M G T I Q E Q T>
 33740 * 33760 * 33780 *
 AAA CAG CTT AAT GAA CAA GTT TTT ATT GAT AAA TTA ACC AAT ATT CCC AAT CGT CGC GCT TTT GAG
 K Q L N E Q V F I D K L T N I P N R R A F E>
 33800 * 33820 * 33840 * 33860 *
 CAG CGA CTT GAA ACC TAT TGC CAA CTG CTA GCC CGG CAA CAA ATT GGC TTT ACT CTC ATC ATT GCC
 Q R L E T Y C Q L L A R Q Q I G F T L I I A>
 33880 * 33900 * 33920 *
 GAT GTG GAT CAT TTT AAA GAG TAC AAC GAT ACT CTT GGG CAC CTT GCT GGG GAT GAA GCA TTA ATA
 D V D H F K E Y N D T L G H L A G D E A L I>
 33940 * 33960 * 33980 *
 AAA GTG GCA CAA ACA CTA TCG CAA CAG TTT TAC CGT GCA GAA GAT ATT TGT GGC CGT TTT GGT GGT
 K V A Q T L S Q Q F Y R A E D I C A R F G G>
 34000 * 34020 * 34040 * 34060 *
 GAA GAA TTT ATT ATG TTA TTT CGA GAC ATA CCT GAT GAG CCC TTG CAG AGA AAG CTC GAT GCG ATG
 E E F I M L F R D I P D E P L Q R K L D A M>
 34080 * 34100 * 34120 *
 CTG CAC TCT TTT GCA GAG CTC AAC CTA CCT CAT CCA AAC TCA TCA ACC CCT AAT TAC GTT ACT GTG
 L H S F A E L N L P H P N S S T A N Y V T V>
 34140 * 34160 * 34180 *
 AGC CTT GGG GTT TGC ACA GTT GTT GCT GTT GAT GAT TTT GAA TTT AAA AGT GAG TCG CAT ATT ATT
 S L G V C T V V A V D D F E F K S E S H I I>
 34200 * 34220 * 34240 *
 GGC AGT CAG GCT GCA TTA ATC GCA GAT AAG GCG CTT TAT CAT GCT AAA GCC TGT GGT CGT AAC CAG
 G S Q A A L I A D K A L Y H A K A C G R N Q>
 34260 * 34280 * 34300 * 34320 *
 TTG TCA AAA ACT ACT ATT ACT GTT GAT GAG ATT GAG CAA TTA GAA GCA AAT AAA ATC GGT CAT CAA

Fig. 4
27/30

L	S	K	T	T	I	T	V	D	E	I	E	Q	L	E	A	N	K	I	G	H	Q>	
34340	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*
GCC TAA ACTCGTTCGA GTACTTTCCC CTAAGTCAGA GCTATTTGCC ACTTCAAGAT GTGGCTACAA GGCTTACTCT																						
A>																						
34420	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
TTCAAAACCT GCATCAATAG AACACAGCAA AATACAATAA TTTAAGTCAA TTTAGCCTAT TAAACAGAGT TAATGACAGC																						
34500	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
TCATGGTCGC AACTTATTAG CTATTTCTAG CAATATAAAA ACTTATCCAT TAGTAGAAC CAATAAAAAA ACTAATATAT																						
34580	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
AAAACATATTT AATCATTATT TTACAGATGA TTAGCTACCA CCCACCTAA GCTGGCTATA TTTCGCACTAG TAAAAATAAA																						
34660	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
CATTAGATCG GGTTCAAGATC AATTTACGAG TCTCGTATAA AATGTACAAT AATTCACTTA ATTTAATACT GCATATTTTT																						
34740	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
ACAAGTAGAG AGCGGTGATG AAACAAAATA CGAAAGGCTT TACATTAATT GAATTAGTCA TCGTGATTAT TATTCTCGGT																						
34820	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
ATACTTGCTG CTGTGGCACT GCCGAAATTC ATCAATGTT AAGATGACGC TAGGATCTCT GCGATGAGCG GTCAGTTTC																						
34900	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
ATCATTGAA AGTGCCGTA AACTATACCA TAGCGGTTGG TTAGCCAAAG GCTACAACAC TGCGGTTGAA AAGCTCTCAG																						
34980	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
GCTTTGGCCA AGGTAATGTT GCATCAAGTG ACACAGGTTT TCCGTACTCA ACATCAGGCA CGAGTACTGA TGTGCATAAA																						
35060	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
GCTTGTGGTG AACTATGGCA TGGCATTACC GATACAGACT TCACAATTGG TGGGGTTAGT GATGGCGATC TAATGACTGC																						
35140	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
AGATGTCGAT ATTGCTTACA CCTATCGTGG TGATATGTCT ATCTATCGCG ATCTGTATTT TATTCAAGCGC TCATTACCTA																						
35220	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
CTAAGGTGAT GAACTACAAA TTAAAGCTG GTGAAATAGA AATTATTGAT GCTTTCTACA ACCCTGACGG CTCAACTGGT																						
35300	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
CAATTACCAT AAATTTGGCG CTTATCTAAG TTGTACTTGC TCTGACCGAC ACAAAATAATG TCGTTCTCA GCATATATCA																						
35380	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
AAATACACAG CAAAAATTG GGGTTAGCTA TATAGCTAAC CCCAAATCAT ATCTAACTTT ACACTGCATC TAATTCCAAA																						
35460	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
CAGTATCCAG CCAAAAGCCT AAACATTG TGACTCAGCG CTAAATATG CGATGCAACA ACAAGTCTT GGATCGCAAT																						
35540	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
ACCTGAGCTA TCAAAATGG TCACCTCATC AGCACTTTGA CGTCCTGTTG CGGACTCGTT TATCACCTGA CCAATCTCAA																						
35620	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
TTATCGGCGT ATTTCTGCTA TGTTGAAACT CACCAATAAC AATAGATTGA GAAGCAAAGT CGAAACAA GCGAGCATGA																						
35700	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
CTATATAGGT CAGTTGGCAA CTCTTGCTTA CCCACTTTAT CAGCGCCCAT TGCAGAAATA TGCGTCCCTG CTTGTACCCA																						
35780	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
CTGCGCTTCA AATAAAGCGC CTTGAGCTGT GGTTGCTGTG ATAATAATAT CTGCTTGTTG ACAAGCAGCT TGTGCATCAC																						
35860	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
AAGCTTCGGC ATTAATGCCT TTTTCTAATA AACGCTTAAC CAAGTTTCA GTTTTGCTAG CACTACGGCC AACTACCAAT																						
35940	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
ACCTTAGTTA ATGAAACGAAC CTTGCTACT GCTAGCACTT CATATTCAAC CTGATGACCG GTACCAAAAA CAGTTAACAC																						
36020	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*	
36040																						
36060																						
36080																						

Fig. 4
28/30

Fig. 4
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37860 * 37880 *
ATCAGTGACA TTAACGTTGC AGCATATTGA AAAGAAACTA TCGATTAGCC TGATC

Fig. 4
30/30

10 20 30 40 50 60
 AATAGATCGACTCGCAAAAGTTGCTTAAGATAGTGTCAATATAGCTTCTTATTTGTAAAT
 70 80 90 * 100 110 120
 ATTGTTTTATGTGTAAACATGTTAGTGTGTAAATGCTGTTAATTATCCTTTGGG
 130 140 150 160 170 180
 ATTGTAATAGCTGATGTTGCTGGCTAATGAGTACTTTAGTTCGGCAATATCTTGCTTTA
 190 200 210 220 230 240
 AATCGCTAACCTCAGTTAATTCAACCCACACTTGTGTATTTAAGGCTCTCTCCC
 250 260 270 280 290 300
 CACCATCGACAAACCAGGATGATGAAACCGGTAAACGTACCAAAGAGACCGACACCTG
 310 320 330 340 350 360
 CAGTCATGAGTAATGCCGCAATGATACGTCCGCCAGTGGTGACGGGTAGTAGTCACCGT
 370 380 390 400 410 420
 AACCAACAGTCGTTATTGTCAAAATGACCACCAAAGTGCCTGATGCCGTTATTGATGT
 430 440 450 460 470 480
 TACTGCCTACTTGATCCTGTTCTAACAAATAAAACCGATAGCACCAAGGTGACAAGGA
 490 500 510 520 530 540
 TGAAGGATATCGCAGATAACCAGCGAAAAGGTGGCTTAAACCGATGTTCAAAATCATT
 550 560 570 580 590 600
 TTAAGATAATTTTGATGAGCGTATATTCTGAATAGATCTTAATACTCTAGCGATACGAA
 610 620 630 640 650 660
 TTATGCGAATAAACTGCAGTTGCTCGACCATCGGAATACTCGACAGTAGGTCAATCCAAC
 670 680 690 700 710 720
 CCCATTCATAAACTGAAATTATTCTCAGCTGGTGAAGCGAATTACAAAGTCAGTGA
 730 740 750 760 770 780
 AAAAGAATAAGCAAATCGTATTATCTACGCTCGTTAATATTCAGTGACGTTACTTGAAA
 790 800 810 820 830 840
 AGGTAAAAATAAGTTGCAGTAGTGTGATACGACCACATGAAGTGATAAAATAAGCATGA
 850 860 870 880 890 900
 AAATCTGAAATGGATTACATCACTGTTGTTGGTGCACCTTTAAGGTTCGTTTCA
 910 920 930 940 950 960
 CAATCTGCTGCCTCGGTTCATGATTTGTTAATATAAACCTTAGTCAGTAGCAAGACAA
 970 980 990 1000 1010 1020
 AATATATTTACATCAATGTATCGTATTATTCAACCGCGCGTGTATTAGACACCAAGA
 1030 1040 1050 1060 1070 1080
 TCGTTGTATATGTTAGTCATGTAGCGATGAGATTATCGCAGACAGGAGAGAATTATGTT
 1090 1100 1110 1120 1130 1140
 TGTTATTATTTACGTACCTAAAGTTAATGTTGAAGAAGTAAACAGGCGTTATTAA

Fig. 5

1150 1160 1170 1180 1190 1200
 CGTCGGAGCTGGCACCATCGGTGATTATGATAGTTGTGCTTGGCAATGTTGGGGACTGG
 1210 1220 1230 1240 1250 1260
 GCAGTTCCAACCTTACTTGGTAGCCAGCCACATATTGGTAAGCTAAATGAGGTTGAATT
 1270 1280 1290 1300 1310 1320
 CGTTGATGAGTTAGAGTAGAAATGGTTGTCGAGCAGAAAATGTAAGGGCAGCAATAAA
 1330 1340 1350 1360 1370 1380
 TGCACATTATTGCTGCGCACCTATGAAGAACCTGCTTATCATATTCTGCAAACATTGAA
 1390 1400 1410 1420 1430 1440
 TCTTGATGAGTTACCTTAAGTTAGATGCACTGCACTTAATTGGTTGCTGTGCTAGGTTA
 1450 1460 1470 1480 1490 1500
 GCAATTAGCAATTGGACCATGTTAGCGATAGTTGGCACAAAGTGATCGATATTAAACT
 1510 1520 1530 1540 1550 1560
 ATCCGATTTCAGATCCCATTAACTGCTGAATTAGGTTCAATTACACTTGTCTAGGGT
 1570 1580 1590 1600 1610 1620
 TTTTCCCGACAGGTGTAACCTGTTACTTGCCTAAGGTTGATAATCTTACCGCATTGGC
 1630 1640 1650 1660 1670 1680
 AGGAGTTACACCTGCACCAGGCATAATACTAATTCTACCATCTGCTTGGTTAACTAACGT
 1690 1700 1710 1720 1730 1740
 TTGGATTAAGGCGCAGCCTCTAGCGCTTGAGCTTGGACAGAGGTTAAACGCTC
 1750 1760 1770 1780 1790 1800
 ACAACCAGCAGTGATCAAGGTCTCCAAGGCTTGTGAGTCAATTACACAGTCGAAAGC
 1810 1820 1830 1840 1850 1860
 GCGGTGGAAGGTTACGCCAGAGTACGTGATGCCACCATAAGCGTTAAAGCTGGCTC
 1870 1880 1890 1900 1910 1920
 GTCAATATTACCATCTGTTAACCGCCAATAACGACCCCTGGACACCGAGTAACCT
 1930 1940 1950 1960 1970 1980
 CATGAATTGATGTCGGAAACCATAATATCAACCTTGTGCTATATACAAAATCACC
 1990 2000 2010 2020 2030 2040
 GGCGCGAGGGCGAATAATGGCATAAAATGGGATCGTGCTAGATCAATAGACTTTGTAC
 2050 2060 2070 2080 2090 2100
 AAAACCTGCGTTGGCGGTCAAGCCACCTAATGCTAATGCCGAGCACAACCTCAATACGATC
 2110 2120 2130 2140 2150 2160
 GGCGCCAGATGCTTGAGCCGTCAAGCAGTGATTCTATATTATCGACACATACTTCTATTGT
 2170 2180 2190 2200 2210 2220
 CATTGTCAATACCTCTTAAAAAGTTATTAAAAATAATAAGCCAGCATAAGTCGT
 2230 2240 2250 2260 2270 2280
 TTTATACAATATGAAAGGGAAAAGGCAGACTAGCTCGCCTAGATCAATTATTATGGCAG

Fig. 5

2290 2300 2310 2320 2330 2340
 AATACTGCCGTATTGTGATTAGAAAAGACAGTTTTAAGCTCAATAGCCGTATCGCGTT
 2350 2360 2370 2380 2390 2400
 GTTATCTACCATCGTGTAACTTTCTGGCCTGGGTGCTTTATTAAACACTGTTTCAGTGGC
 2410 2420 2430 2440 2450 2460
 TGGATTAGGGTGAAATGATTCTTTTCAAATCTGTTTTGTATTTGAACGTACCTGT
 2470 2480 2490 2500 2510 2520
 AATGTCTTGCTGCTCACGAAGACGTACAAATATTGGTTGCGCATAGCTGGTAGTGCCGC
 2530 2540 2550 2560 2570 2580
 ATTGACATGTTGATAGAATTCAAGACGCTGAAAATTCAATGAATAGGGCAATTCAAAGTCAG
 2590 2600 2610 2620 2630 2640
 CGCGACCATGCCTGCTCGGCCATCGTGTGAGCTGGGAGCTTGACACCATAAGCCACACTTG
 2650 2660 2670 2680 2690 2700
 CTCAATTCGCACAAAATCGTTAACTTGAGCTCTACTTGCCTCGTGGCGACATTTCAACC
 2710 2720 2730 2740 2750 2760
 TTTCCAGCGGAATGTATCACCTAATCTATCCACAAAGGAAATATGGCGATAACCTTGGTA
 2770 2780 2790 2800 2810 2820
 ATGAACGAGATGCCGGTATTAAAATAACAGTCACCGTCTTTAATACTGACTTAAATAG
 2830 2840 2850 2860 2870 2880
 CTTTTTATTACTTCGTTGTATCGGTATAACCATAAAATGGTGAACGTTAGTTATCTT
 2890 2900 2910 2920 2930 2940
 TGTTAGCAGTAGCCCTGTTCTCCGTTTACTTGGTCATTTCCTTCGCATTATA
 2950 2960 2970 2980 2990 3000
 CACAGGTTTGTCAATTGCAATATCATATTGTATGACGGTAAAGCAAGTGGAGTAACCCC
 3010 3020 3030 3040 3050 3060
 CGCTGTATGCCGTAAAGTTCAGCGCATTGGAGAACACAAGATTACACTCACTGGCGCCATA
 3070 3080 3090 3100 3110 3120
 GAATTCAATTATGCTCGATCCAAAACGTTGGAAATGATCCAAATTTCGGGGCG
 3130 3140 3150 3160 3170 3180
 TAATCCATTACCTATGATTTCTTATATTATGCTGTTGTCTTATTGCTAGGCGGTAC
 3190 3200 3210 3220 3230 3240
 ATTTAATAAAACGGCAGAGCTGCCGATGTAAGTAAACGCAGTGGCATTATGAGCACG
 3250 3260 3270 3280 3290 3300
 AACTTCATCCAAAAGCGACTTGAACTGAATTTCAGAAAGTGCAGGGTTGCTGCGCT
 3310 3320 3330 3340 3350 3360
 ACCAAACACGGCGCTTAATGACACTGTCAGTCAGTGCATTGTTATGGTATAGGGGGAGTGATAA
 3370 3380 3390 3400 3410 3420
 ATACAATACATCATCAGCTGTTAAGCGTAATGATGCCATCCCCATGCCTGCCATGGATT

Fig. 5

3430 3440 3450 3460 3470 3480
 AAACCAACGGTGATGGCTCATTCTGCTGCTTTGGCAGTCCAGTTTCCCGAGGTAAA
 3490 3500 3510 3520 3530 3540
 GATATAAAACGCGCAATGCTTAAGCTGTATTTGTGCTGTTGATTCAAGGGTTCAAACTGA
 3550 3560 3570 3580 3590 3600
 ATATCCTGCGACTAGTGTAGATAATGTTTATAACCACACTCATGTCTGGCGTTCTAA
 3610 3620 3630 3640 3650 3660
 AGCGGGTACGTAAAAGACATTCTGTTGAATGTCGATGACAAATTGGTTCAATATTATT
 3670 3680 3690 3700 3710 3720
 AATGGCGGATGTGTATAGTCATCTGCGATGAGTAATTGGTATCGACCACGCTAAGACT
 3730 3740 3750 3760 3770 3780
 ATGTTCGAGGATTGAATCCCCTGTCGTATTTATCATAACAAGCAATCGCGCCAAGCTT
 3790 3800 3810 3820 3830 3840
 GACAACGTGCGAGGGCAATAATGATGGTTCAAGGCCTGTTATCGAGCATGATGGCGACTTT
 3850 3860 3870 3880 3890 3900
 ATCATTTCACCAATGCCGTATTCATGAAGGAAATGGCATATTGATTGCTTGCTTATT
 3910 3920 3930 3940 3950 3960
 CAATGAATCGTAACATAACGCTGGTCTTAAATTGTATTGCGATCAAGTCAGAGTTATT
 3970 3980 3990 4000 4010 4020
 GACAGCTTGCTGCTCTAGTAATAAACCAATAGACATAAACGTTCGGGCTTGCTTGTG
 4030 4040 4050 4060 4070 4080
 TAAGTGCCATAAGCCTTGATGATTGGCTTGGGTTTTAATAGATTGATGGTACTTTT
 4090 4100 4110 4120 4130 4140
 CAGGAATTGTTGCCGGTTATAACAGTCATAAGCTAATTCTTTATCAAGAAGAGGGT
 4150 4160 4170 4180 4190 4200
 TATGACACCAAATAATGGGTACCGCTGGTTAATTGGTAGACTAAATGTGTTGTT
 4210 4220 4230 4240 4250 4260
 TTGCTGTGATAATGCGACGTTCAAACAAACTTGAGAAGGTAAAAAAATAGCATTAA
 4270 4280 4290 4300 4310 4320
 TTGAACATCAATACTAATGTGTTGAATATCAATCAAGTTCTAACTGTGCGAGCACCG
 4330 4340 4350 4360 4370 4380
 TGCTTTAGCAAACATGCCATGTGCTATTGCTGTTAAACCCATTAGTTCGCTGGGAT
 4390 4400 4410 4420 4430 4440
 AAAATGTAAATGGATTGGATTGTCCTTGGAGATATAAGCATATTATACGTCAA
 4450 4460 4470 4480 4490 4500
 AGGACTAAATTAAACAATGAAATCGGCTCGTAAGCATAATTGCTGGCGTATTTACTAT
 4510 4520 4530 4540 4550 4560
 TTTCTCACCGCTGGAACGTTGAGATCGTGGCACGTTTCGCTGTTCGTTCTGTAA

Fig. 5

4570 4580 4590 4600 4610 4620
 GAATGTCGATGTACACTCCCACGCAAATTGTCCATCTACAAACACATCAATATGAGTATC

 4630 4640 4650 * 4660 4670 4680
 AATGAAACGTCCTGTATCCGTTATGTACTCCTTAATTACACGACATGTGCTCGTCAATAT

 4690 4700 4710 4720 4730 4740
 CGCGTTAATGCTATCGGTTGATGTTGTATGCGATTCGATAATGGACTAGTCCTAA

 4750 4760 4770 4780 4790 4800
 TATAGATATCGGAAATTGTGTTGATGTCATGAGTTTCATCAATAATGGAAAGATCATCAC

 4810 4820 4830 4840 4850 4860
 AAATGGATAAGTAACCGGTACATAGTTGTGTTATTAAACCCACAGCATTAAATATATTG

 4870 4880 4890 4900 4910 4920
 CTTTAAATTCGCTGATCTATTGGTCACTGATACTAAATTGCTCAGTACACACTTG

 4930 4940 4950 4960 4970 4980
 TGTCGACCAAGTGTTCATCAGTGTAAACAAATTGATTGACCACTGCTTCACATATAA

 4990 5000 5010 5020 5030 5040
 AAGCGAGATAATCGGTTGCTTGTTAACAGTGTGATCTGGTTAGCGTGCATTGAAATAAT

 5050 5060 5070 5080 5090 5100
 TCATATAAGAGTATGTAGCATTATGTTAATATTTGTTGGAAAGTTGAATTGGCGAAT

 5110 5120 5130 5140 5150 5160
 CCGTAATCGGTTATGGCAGTCGGTCAAATACTTCAGGTAAACTCGTTACTCATACCAT

 5170 5180 5190 5200 5210 5220
 TGATAGTGTAAAGTGTGATTGACTGAATAAGAATAGAGCTAAAGTGGAAAAATTATGCA

 5230 5240 5250 5260 5270 5280
 AGATGCGGGTATGTTATTACGCATTGCTTATGAGGCAATGAAAGAGTTAGAGGTTGATGT

 5290 5300 5310 5320 5330 5340
 CATTGAAGTACTTCTCGTTGTAACATAAGTGAAGAAGTACTGAATGATAAGGATCTCG

 5350 5360 5370 5380 5390 5400
 CACACCTAATCATGCACAAACACATTGGCAAGTATTAGAAGACATATCACAAGATCC

 5410 5420 5430 5440 5450 5460
 TAACATCGGCATTCACCTGGTGAGAGAATGCCAGTGTACGGGGCAGGTATTACAGTA

 5470 5480 5490 5500 5510 5520
 TCTTTCTCAGTAGTCCTACATTGGTACTGGCTGGAACGCGAACAAAATACTTCG

 5530 5540 5550 5560 5570 5580
 ATTAATCAGTGATGCGGCGAGTGTCTATCAAGATGGAAGGCTGTGAAGCGCGATTATC

 5590 5600 5610 5620 5630 5640
 TGTGAACCTAGATGGTTAGCGGAAGATGCGAATCGTCATTGAATGATTGCCTAGTGAT

 5650 5660 5670 5680 5690 5700
 CGGTGCATTAAATTGGTTATATGTGACAGAAGGCGAATTAAAGTAAGCAAAATAGC

Fig. 5

5710 5720 5730 5740 5750 5760
 CTTTGCTCATGCTCGCCCGAAAGATAATTACTGCCTATACCAATGTATTTACATGTCCGAT
 5770 5780 5790 5800 5810 5820
 TGAGTTTGCTGCCGAAGATAATTATATTTATTCGATGCTGATTACTCGAACGTCCTTC
 5830 5840 5850 5860 5870 5880
 TTCGCATGCGGAGCCTGAGCTATCGCCTTACACGATCAGCTTGAAGCCGTAAGCGTAAAGC
 5890 5900 5910 5920 5930 5940
 CAAGTTAGAACTGCAAGATTTAGTGGATAAAGTACGTAAGGTTATTGCACAAACAAC TTGA
 5950 5960 5970 5980 5990 6000
 GTCTGGTGTGGTGACTTTAGAAAGTATCGCCACTGAACCTGACATGAAACACGTATGCT
 6010 6020 6030 6040 6050 6060
 AAGAGCGAAGTTAGCTGACATTGATTATAACTTTAACAAACTCGCTGATTTCGTTG
 6070 6080 6090 6100 6110 6120
 CGAGTTATCAAAAAACTGTTGGCGAATA CGGACGAGTCTATTGATCAGATTGTCTATCT
 6130 6140 6150 6160 6170 6180
 CACTGGTTTTCTGAACCAAGTACTTTTATCGTGCCTTAAGCGCTGGGTTAAATGAC
 6190 6200 6210 6220 6230 6240
 GCCAATTGAATATCGCCGTAGCAAACCTCGCGGTTAGGCATGCTAATCAACACGAGTCCTA
 6250 6260 6270 6280 6290 6300
 AAAATTGCTGCTTAGTGCATAGTGCATAGTGCATAGTGCTAGTAAGCCAAGTACAAAGC
 6310 6320 6330 6340 6350 6360
 GTTAAAGTTAAGTACTTGAGCGAACCATCAGACACCACCTACTAGATTAAAGCACCTATTA
 6370 6380 6390 6400 6410 6420
 ATGATTGACCACAAATTCTGATCGTATTGCCTGTGATCCCTGCAGCTTGAGGTTGCGCAA
 6430 6440 6450 6460 6470 6480
 AAAAAGCTATCGCTTCAGAACATCAACTGGCTTACCACTTGTAAATGAATTACATAC
 6490 6500 6510 6520 6530 6540
 GACGACCAGCTTCACGAACTGTAAATGGAATCGCTGCTGTCATTTGTTCAATAAAGC
 6550 6560 6570 6580 6590 6600
 CTGGTGCAACAGCATTAATGGTGATGTATTTGTCATGCAAGCGGAGTTGCATTGCA
 6610 6620 6630 6640 6650 6660
 CATAACCAATGACTGCGGCCTTAGACGTTGCATAATTAGTCTGACCAAAGTTACCGCAA
 6670 6680 6690 6700 6710 6720
 TCCCACACTCGAACAGACACACAAACATGCGGCCATAGTCGTTGAGCAGATCATCATT
 6730 6740 6750 6760 6770 6780
 GCAGTCGCTCATTGATTCTTCCATTGCCGACAAGTTAATATCCATCAGTACATCCCAAT
 6790 6800 6810 6820 6830 6840
 GGTTATCCGGCATACTGCTAGCGTTGTCTTTGTTACCCCGCATTATGGACGATGA

Fig. 5

6850 6860 6870 6880 6890 6900
 TATCAAGCGACTGTTCTCGCACAAAGTCAGCAATGATATTGGGGCGTCAGCAGCGGTAA

 6910 6920 6930 * 6940 6950 6960
 TATCAGCAACAATGCTGCTACCTTCAGCAATGAGCTACTTTCAAGGTCCTGTTTA

 6970 6980 6990 7000 7010 7020
 ATGCCGGAATGTCTAACGAAATAACATGTGCGCCATCACGGGCGAGTGTTTCAGCAATAG

 7030 7040 7050 7060 7070 7080
 CAGCCCCGATGCCACGTGATGCACCAAGTGACAAGTGCTGTCTTCCTTGTAAATGGTTTG

 7090 7100 7110 7120 7130 7140
 CCGTGTACTTGTTCTGTTAAATAACTCGTTAATAACTCGTTAATAACTCGTTAATAG

 7150 7160 7170 7180 7190 7200
 CCCCATTAAATCGAACCGGGTTTACGTTAATAACCTGTGCTGAGATATAGGCTGATTTG

 7210 7220 7230 7240 7250 7260
 CTGAGGTTAACGAAACGTAGCGGGGCCTCTAATAATTGCTCACTACCAGGTTGTACATAGA

 7270 7280 7290 7300 7310 7320
 TAAGTTGACAGGTACTACCATTCTGCCTATTCTTGGCGACACTGCGACAAAACCCCTT

 7330 7340 7350 7360 7370 7380
 CTAAAGATCTTGTACAGTCGCGTAGCTTACATCGTCAAGATGTTCACTCGGATGACCTA

 7390 7400 7410 7420 7430 7440
 ACACGATCACTCTGCTGCATGGCGAGAGCTGCTTAATTACAGGTTGAAAAAAACGATGTA

 7450 7460 7470 7480 7490 7500
 ATGCACTTAATTGCTTGCTGTTCTTAATGCCCTGAGGCGTCGAAGATAATTACCGTTGAAGC

 7510 7520 7530 7540 7550 7560
 GATCTGTTTAGCGATAGCATTAAAGGCTAATAGGTGTCGCGACTAAAGACGTTGATTAA

 7570 7580 7590 7600 7610 7620
 ATTCAATATTAAAGATCGGCTAACGCTGACGTGTTATTAGGATAAGAAATCGTGACTTCAG

 7630 7640 7650 7660 7670 7680
 CATCTTAAATGTGTTAAGAATGGGTTAATTAAATTGCTGTTGCTGGCTGCGCCGATGA

 7690 7700 7710 7720 7730 7740
 GTAAGTTGCCAGAGATGAGATCGGTTCCCTGATCGTAGCGTGTAAACGTAACCGGTCGT

 7750 7760 7770 7780 7790 7800
 GCAGATTAAGCGCTTAAATAAACCTGATGTCCACTGCCATTAGCGAGTTTGCATG

 7810 7820 7830 7840 7850 7860
 TATCCGTCACTTCTAATCCTGTTAGTGAACAGTTGAATCTCGAAGATGTACATGT

 7870 7880 7890 7900 7910 7920
 GTTAAAAAATTATCTGATAGCTATGACTTATCTGCCACTACGTAATAATAATAGACCAAGT

 7930 7940 7950 7960 7970 7980
 TCATTACATCGTTAATCGATATAGTATAACTAAACTAAAGTAAATTATAATGATAAGAC

Fig. 5

M

7990 8000 8010 8020 8030 8040
 TGTTATCGTACTCGGATCAAACCTCTGATCAGCAAATAATCAAATTAGAGTTTTATTTA

 8050 8060 8070 8080 8090 8100
 AACTTGTATCAACAAATGTTACATTAATGTATCTTACGTCTAATGTGCTACGGGCATATT

 8110 8120 8130 8140 8150 8160
 AAGTCACTAAATTAAAGGAATAAACCATGACAGGTCAAACAATAAGAAGAGTAGCAATT

 8170 8180 8190 8200 8210 8220
 TCGGCAGGTAAACCGTATCCCCTTGCACGTTCAAATACAGCGTATTCAAACACTAAGTAACC

 8230 8240 8250 8260 8270 8280
 AAGATATGCTGACGGAAACTATCCGTGGCTGGTAAATATAACCTACGTGGTGAAC

 8290 8300 8310 8320 8330 8340
 AACTGGGGGAAGTTGTTGCTGGTGCAGGTAAATTAAGCATTCTCGTGAATTAAACAC

 8350 8360 8370 8380 8390 8400
 GTGAAGCCGTGCTAAGTGCAGGTCTTGCACCTGAAACGCCTTGTATGACATTCAACAAG

 8410 8420 8430 8440 8450 8460
 CTTGTGGTACTGGTCTAGCTGCAGCTATCCAAGTAGCAAACAAAATTGCGCTTGGTCAA

 8470 8480 8490 8500 8510 8520
 TAGAAGCGGGTATTGCTGGTGGTCTGATACGACATCAGATGCACCGATTGCAGTCAGTG

 8530 8540 8550 8560 8570 8580
 AAGGCATGCGTAGTGTATTACTTGAGCTTAATCGAGCTAAACGGGTAAGCAACGTTGA

 8590 8600 8610 8620 8630 8640
 AAGCACTATCTCGTCTACGTCTAAACACTTTGCGCCACTAACGCCCTGCAAATAAGAGC

 8650 8660 8670 8680 8690 8700
 CGCGTACCAAAATGGCGATGGCGATCATTGTCAAGTAACAGCGAAAGAGTGGAAATATCT

 8710 8720 8730 8740 8750 8760
 CACGTGAAGCACAAGATGCATTGGCCTGCGCAAGTCATCAAAATTAGCTGCAGCATATG

 8770 8780 8790 8800 8810 8820
 AAGAAGGTTCTTGATACGTTAGTTCACCTATGCCGGCTAACGAAAGATAACGTAT

 8830 8840 8850 8860 8870 8880
 TACGCGCAGATAACACAGTTGAGAACTGGCTAAATTGAAACCTTGTGTTGATAAAAGTAA

 8890 8900 8910 8920 8930 8940
 ACGGCACATGACGGCGGTAAACAGTACTAACCTTACCGATGGAGCATCAGCTGTATTAC

 8950 8960 8970 8980 8990 9000
 TTGCAAGTGAAGAATGGGCAGCGGCACATAACTTACCGTACAAGCTTACACATTG

 9010 9020 9030 9040 9050 9060
 GTGAAACGGCCGCTATCGACTTCGTTGATAAGAAAGAAGGTCTGTTAATGGCGCCTGCAT

 9070 9080 9090 9100 9110 9120
 ACGCAGTGCCAAAATGTTGAAGCGTGGQCTTACATTACAAGACTTCGATTACTATG

Fig. 5

9130 9140 9150 9160 9170 9180
 AAATACATGAAGCATTGCTGCGCAGTTATTAGCAACGCTAGCAGCTGGGAAGACGAAA

 9190 9200 9210 9220 9230 9240
 AATTCTGTAAAGAAAAACTGGGTCTAGATGCTGCGCTGGTTCAATTGATATGACCAAGT

 9250 9260 9270 9280 9290 9300
 TAAACGTAAAGGGAGTAGCTTAGCCACGGTCACCCATTGCCGCAACTGGTGGTCGTG

 9310 9320 9330 9340 9350 9360
 TTGTCGCTACGCTAGCGCAATTACTGATCAGAAAGGTTAGGTCTGGTTGATCTCGA

 9370 9380 9390 9400 9410 9420
 TTTGTGCTGCTGGTGGTCAAGGTATCACGGCAATTAGAGAAATAACGCACGTGTTAT

 9430 9440 9450 9460 9470 9480
 TATCTATTGATTAAGCTGTCCTGAGATACTGGATATTTAAATAACGCCAATACTGC

 9490 9500 9510 9520 9530 9540
 AGAGTATTGGCGTTTTGTAAATCCAATTCTATATAACGGTGCATTTAAACACTTA

 9550 9560 9570 9580 9590 9600
 ATTTCCGGCATTGGTATCATAAAAAGCAGCACCGAAGTGCTGCTTGATTGTAGATTAAC

 9610 9620 9630 9640 9650 9660
 CTATTAAAATAGAGAGGCTAGAATTAGTCTCGTATGCTTCATTATGTACGCCAGCTGCA

 9670 9680 9690 9700 9710 9720
 CGACCCGATGGATCAGCATTGTTGGAAACTTCATCCCAAGCTAATGCTCTACAGTT

 9730 9740 9750 9760 9770 9780
 GAACAAGCAACGGATTACCAACGGTACCGCATTGCTGCTGAATCACCTGGGAAGTGA

 9790 9800 9810 9820 9830 9840
 TCTTCAAAGATGGCACGATAGTAGTAACCTTCTTCGTATCTGGTGTGTTATTGGGAAC

 9850 9860 9870 9880 9890 9900
 TTAAATGCTGCACTTGCTAACATTGATCAGTTACCGCTTCTCAACGTGTACTTTAAGT

 9910 9920 9930 9940 9950 9960
 TGGTCAATCCAAGAATAACCAACACCATCAGAGAATTGTTCTTTGACGCCATACAATT

 9970 9980 9990 10000 10010 10020
 TCTTCAGGTAGTAAATCTTCAAATGCTTCGAATGATGTTTCTCAATGCGGTGCC

 10030 10040 10050 10060 10070 10080
 GTGATCATTTAGTTCAAGGGTTAGACGCATTGACGCATCAACAAATTCTTATCTAAG

 10090 10100 10110 10120 10130 10140
 AAAGGAACACGTGCTTCGATGCCCAAGCTGCCATAGATTGTTGCACGTAAAGCAATCA

 10150 10160 10170 10180 10190 10200
 AACATATGTAATTATTTACTTACGTACCGTCTTCAATGGAATTCTTCGCATTGGC

 10210 10220 10230 10240 10250 10260
 GCTTGTGGAAGTACAAGTAACCACCGAACAGTTACAGCACCTCACCAAGAAAGCACC

Fig. 5

10270 10280 10290 10300 10310 10320
 ATCTTAATCCCCATGGCTTAATTTACGTGCCATTAGGTACATAGGGGTTGATGCACGA

 10330 10340 10350 10360 10370 10380
 ATTGTTGTTACATCGTAGGTTCAATGTGGTAAATCACGTGCGTAAAGCGTCGATACCT

 10390 10400 10410 10420 10430 10440
 TCTTGCACAGTAAATTCAATTGAATGATGGATAGTACCTAAGTGATCTGCCACTTTTGT

 10450 10460 10470 10480 10490 10500
 GCAGCGGCTAAATCTGGAGAACCATTTAGGCCTACAGAGAAAGAGTGTAGTTGTGCCAC

 10510 10520 10530 10540 10550 10560
 CATGCTTCGGTTTACCAACCGTCTCAATACGACGTTGCATACTGTTGGGTGATTGCT

 10570 10580 10590 10600 10610 10620
 GAAATAACAGATGAATCTAACCGCCTGATAATAACGCGTAAGGTACATCACACATT

 10630 10640 10650 10660 10670 10680
 AATTGACGTTAACTGCATCTCCAAACCTGCTTAACAAACGCTTTATCACCACCATT

 10690 10700 10710 10720 10730 10740
 TGTGCAACGTTATCAAAATCTTCCAATCACGTTGATAATAAGGCGTGACTACACCATTCC

 10750 10760 10770 10780 10790 10800
 TTACTCCACAGGTAATGACCTGCTGGGAATTCTCAATTGAGTACAAATTGGCACTAGT

 10810 10820 10830 10840 10850 10860
 GCTTTCATTCAGAGGCAACATAAAAGTTACCGTGTTCATCATAGCCCCGTATAAAGAGGG

 10870 10880 10890 10900 10910 10920
 ATGATACCGATATGGTCACGGCCAATCAGGTAAGCGTCTCTGTTCGTCATATAAAGCG

 10930 10940 10950 10960 10970 10980
 AAAGCAAAATACCATTAGATCATCTAAAAATTGTGTGCCTTTCTTATATAGCGCA

 10990 11000 11010 11020 11030 11040
 AGTATCACTTCGCAATCTGATTCTGTTGGAATTCAAAGTCTACGTTCAGCGTTCTT

 11050 11060 11070 11080 11090 11100
 AAATCTTGTGGTTATAAATTCAACGAACTACAGCAAGTACGTGTCTTTCTTCATTA

 11110 11120 11130 11140 11150 11160
 TATAGCGGCTGTGCACCATTATTACATCGACAATAGCAAGACGTTCATGAACATAAAATA

 11170 11180 11190 11200 11210 11220
 GCATTGTCACTTGTATAGATACCTGACCAATCTGGGCCGCGGTGACGTAGTAACCTTGAT

 11230 11240 11250 11260 11270 11280
 AGTTCTAGTGCTTGTGCGAAGAGGTTAATGTCTGATTTGATGTCTAGAATTCCGAAT

 11290 11300 11310 11320 11330 11340
 ATTGAGCACATAACTAATTCCCTCTGGGGCTGCGTCTGCAGCTAACTTCTAAATAGTGT

 11350 11360 11370 11380 11390 11400
 GTCTAATTGCCACATTGTAGATTTAATGCAAACATTAATGATAAAACATTATAAAAAA

Fig. 5

11410 11420 11430 11440 11450 11460
 TGTAATTCAATGTGGAATCGATAATTTAATGGCTTAAAGTGAAGATCCATTAAATTGTGA
 11470 11480 11490 * 11500 11510 11520
 TGGCGAGGTGATAGACCAATGTAGACCTTAATGAATAAAGCAGGCACGATTGAATCCATT
 11530 11540 11550 11560 11570 11580
 CAACGCAAAGTGGTACTAACTATTGTTTAAACGTTATAAATAGTGTAAAGGTTATA
 11590 11600 11610 11620 11630 11640
 AGTAAATAATTAAAAACAATAATAATCCACATGCATTAAATTATCATGATAAAACCGCT
 11650 11660 11670 11680 11690 11700
 ATATCTCAATGGCAATTGGGATAAGTGTAAAATATATGTAAAATGAATGAGTTGACTTG
 11710 11720 11730 11740 11750 11760
 CTTTTTTACACTAAGTGTGAAATTAAAGCTAGATGTCGTTAGCATTGATTAATAA
 11770 11780 11790 11800 11810 11820
 CGTACTAAAATACGACATCTAGTATAGAAATTAAACAGTTGGTTTGATAGCATAA
 11830 11840 11850 11860 11870 11880
 CTGCATAAAACTAATCAGCTATTGTCGTAAATTTGTAAATTAAATAGGTTAATAA
 11890 11900 11910 11920 11930 11940
 AATTATATGTCTGATAAAATATAACCGTACGACCTTCCTTAAAGACGTTTGCTG
 11950 11960 11970 11980 11990 12000
 CCTAAGTTTGGCCTGTGTGGTTCGGGTGTTGCAATATACTTATTAGCTTTATGCCA
 12010 12020 12030 12040 12050 12060
 GTAAAGCCCGTGATAAATTGCTCGATTCAAGCGAAATTGTTAGTCTAAATG
 12070 12080 12090 12100 12110 12120
 ATGGCAAAGCGTAAAAGGTAGCAAAGATCAATTATCTATGTGCTCCCTGAAATGGAT
 12130 12140 12150 12160 12170 12180
 GATACGGAACAAGACCGTATAATCATGGTCAATCTAGTTACTTTGTCAAACATCTTA
 12190 12200 12210 12220 12230 12240
 AGTTATGCAGAGCCAAGTGCAGCTAGTCGTGCTTATAACCGTGACCGTATGATAGTCAT
 12250 12260 12270 12280 12290 12300
 GGTGGCGAGAATTATTCCGCTACTTGAACAAGGTAAGGCTTGTATCTTATTAGTGCAG
 12310 12320 12330 12340 12350 12360
 CATAGCTTCGCTATTGATTTCGAGGTTACACATTGCTTCTTATGGCGGCCATTG
 12370 12380 12390 12400 12410 12420
 ACTATGTTAACATTGAGAATGAGTTGTCGATTGGCTGATGACACGTCAACCGC
 12430 12440 12450 12460 12470 12480
 ATGTTGGAGGCAGTGTTCACCGCAAGGCAGGGCTAGGGCTCTAGTTAAATCACTT
 12490 12500 12510 12520 12530 12540
 AAGAGCGGTGAAAGCTGTTATTACTTACCTGATGAAGACCATTGGACCTAACGCTAGTGTA

Fig. 5

12550 12560 12570 12580 12590 12600
 TTTGCGCCTTATTCGACTCAAAAGCAACTTACCTGTAATGGGCAAGCTAGCAGAA

 12610 12620 12630 12640 12650 12660
 AAAACAAATGCACTCGTTGTTCTGCGCATATAATGAATCACTAGGTAAATT

 12670 12680 12690 12700 12710 12720
 GAAACCTTATTGACCAGCAATGCAAAACTTCCATCAGAAAGCCCAGAACAGATGCA

 12730 12740 12750 12760 12770 12780
 GTGATGATGAATAAGAGATTGAAGCCTGATTGAATGTGGTGTGATCAATATATGTGG

 12790 12800 12810 12820 12830 12840
 ACACCTAGATTATTGAGAACACGTCCGGACGGTAAAAAAACTACTAATAAGTTAATA

 12850 12860 12870 12880 12890 12900
 AACACCATAATCTCGTTGAATATGGTGTACCCCCCTGAATACCCCTCTAAATTAA

 12910 12920 12930 12940 12950 12960
 CAAAAAAAGCCATTACGTAACATCTAATGATGATTAGCCTGCACTGCTTGTGTTTA

 12970 12980 12990 13000 13010 13020
 GTCTTAAGAGCCTAATAAACTTGATCTAGGTATAGATTCTGTCTTACGTAACGCG

 13030 13040 13050 13060 13070 13080
 ATCTATTTTTTAACCGATAAGTTGTTATAATTAGTTCATATGAAAGAGATATCGTTTC

 13090 13100 13110 13120 13130 13140
 AGTAAAGCTATTCGTTCAATAGATAATTATAGTCATATTCGTAATGACA

 13150 13160 13170 13180 13190 13200
 ATCATTTCATCTAGACTATAGATAAGAACGAAATTAGTAAGAACATTAAATTTC

 13210 13220 13230 13240 13250 13260
 AAGAATATAAAATATCCCATCGGAGCTATAAGAATGAAAAAGACTAAAATTGTTGTACA

 13270 13280 13290 13300 13310 13320
 ATTGGTCCAAAAACTGAATCAGTAGAGAAACTAACAGAGCTTGTAAATGCAGGCATGAAC

 13330 13340 13350 13360 13370 13380
 GTTATGCGTTAAATTCTCATGGTAACTTGCTGAACATTAGTGCCTATTCAAAAT

 13390 13400 13410 13420 13430 13440
 ATCCGTCAAGTAAGTGAACCTGAATAAGAAAATTGCTGTTTACTGGATACTAAAGGT

 13450 13460 13470 13480 13490 13500
 CCAGAAATCCGTACGATTAAACTAGAAAACGGTGACGATGTAATGTTGACCGCTGGTCAG

 13510 13520 13530 13540 13550 13560
 TCATTACGTTACAACAGACATTAACGTGGTAGGTAATAAGACTGTGTTGCTGTAACA

 13570 13580 13590 13600 13610 13620
 TATGCTGGTTTGCTAAAGACCTTAATCCTGGTCAATCATCCTGTTGATGATGGTTA

 13630 13640 13650 13660 13670 13680
 ATTGAAATGGAAGTTGTTGCAACAACTGACACTGAAGTTAAATGTACAGTATTAAATACT

Fig. 5

13690 13700 13710 13720 13730 13740
 GGTGCACTTGGTAAAAATAAAGGCGTTAACCTAACATCAGTGTAGGTCTACCTGCA
 13750 13760 13770 13780 13790 13800
 TTGTCAGAAAAAGATAAAGCTGATTAGCGTTGGTTGTGAGCAAGAAGTTGATTTGTT
 13810 13820 13830 13840 13850 13860
 GCTGCATCATTATTACGTAAGGCTGATGATGTAAGAGAAATTCTGAAATCCTATTTAAT
 13870 13880 13890 13900 13910 13920
 AATGGTGGCGAAAACATTCAAGATTCTGAAAATTGAAAACCAAGAAGGTGTAGACAAT
 13930 13940 13950 13960 13970 13980
 TTTCGATGAAATCTTAGCTGAATCAGACGGTATCATGGTTGCTCGTGGCGATCTCGGTGTT
 13990 14000 14010 14020 14030 14040
 GAGATCCCAGTTGAAGAAGTGTACATGGCACAGAACAGATGATGATCAAAAAATGTAATAAA
 14050 14060 14070 14080 14090 14100
 GCAGGTAAAGTTGTAATTACTGCAACACAAATGCTTGATTCAATGATCAGTAACCCACGT
 14110 14120 14130 14140 14150 14160
 CCAACACGTGCAGAACGGCGATGTTGCCAATGCTGTGCTTGACGGTACCGACGCGGTA
 14170 14180 14190 14200 14210 14220
 ATGCTTCTGGTGAACACTGCGAAAGGTAAATACCCAGTTGAAGCTGTCTATCATGGCA
 14230 14240 14250 14260 14270 14280
 AACATCTGTGAACGTACTGATAACTCAATGTCTCGGATTAGGTGCGAACATTGTTGCT
 14290 14300 14310 14320 14330 14340
 AAAAGCATGCGCATTACAGAACGCTGTGTAAAGGTGCGGTAGAACACAGAAAAATTG
 14350 14360 14370 14380 14390 14400
 TGTGCTCCACTTATTGTTGCAACTCGTGGCGTAAATCAGCAAAATCTGTTCGTAAA
 14410 14420 14430 14440 14450 14460
 TACTTCCGAAAGCAAATATTCTTGCTATCACAAACAAATGAAAAAGCAGCGAACAGTTA
 14470 14480 14490 14500 14510 14520
 TGCCTAACTAAAGGCGTAAGCAGCTGCATCGTTGAGCAGATTGATAGCACTGATGAGTTC
 14530 14540 14550 14560 14570 14580
 TACCGTAAAGGTAAAGAGCTTGCATTAGCAACTGGTTAGCTAAAGAAGGCGATATCGTT
 14590 14600 14610 14620 14630 14640
 GTTATGGTATCAGGTGCGTTAGTACCATCAGGTACAACGAATAACGGCATCTGTTACCAA
 14650 14660 14670 14680 14690 14700
 CTTTAAGTTGCCATATTGATATTATAAAAAAGAGAGCGTATGCTCTTTTTTATATCT
 14710 14720 14730 14740 14750 14760
 GTAGTTATATGTCGTCACAAAAAAATGATAAAGAGTACATAAACTATTAATATAGCGTA
 14770 14780 14790 14800 14810 14820
 ATATATAATGATTAACGGTGATGAAAGGGTTAAATAAATGGATAGTGCTAAACATAAAAT

Fig. 5

14830 14840 14850 14860 14870 14880
 TGGCTTAGTCCTTCTGGCGGTGGTGCAGAAAGGTATTGCTCATCTGGTGTATTAAAATA

 14890 14900 14910 14920 14930 14940
 CCTGTTAGAGCAAGATATAAGACCGAATGTAATTGCGGGTACAAGTGCCTGGCTATGGT

 14950 14960 14970 14980 14990 15000
 TGGTGCACCTTATTGCTCAGGACTTGAGATTGACATTACAATTCTCATCGATGT

 15010 15020 15030 15040 15050 15060
 AAAACCTTTCTTGGAAAGTTACCCGTGCCCGTGGCTTATAGACCCGGAAAATT

 15070 15080 15090 15100 15110 15120
 ATATCCTGAAGTGCTAAAATATATCCCCGAGGGATAGCTTGAGTACCTAACCTGAATT

 15130 15140 15150 15160 15170 15180
 GCGCATTGTTGCCACCAACATGTTACTCGTAAAGAGCATATATTAAAGATGGCTCCGT

 15190 15200 15210 15220 15230 15240
 GATTAATGCCTTATTAGCATCAGCCAGCTACCCCTTAGTTTTCTCCGATGATCATTGA

 15250 15260 15270 15280 15290 15300
 CGATCAAGTGTATTAGATGGCGGTATTGTTAACATTCCCCGTGAGTGTCAATTGAAGA

 15310 15320 15330 15340 15350 15360
 TGATTGCGATAAAATAATCGGCGTATACGTGTCGCCATTGTCAGGTCGAAGCTGACGA

 15370 15380 15390 15400 15410 15420
 ACTCTCGAGTATAAAAGACGTGGTATTACGTGCGTTACGCTGCAGGGTAGTGGTGCTGA

 15430 15440 15450 15460 15470 15480
 ATTAGATAAAACTATCGCAATGTGATGTGCAAATTATCCAGAAGCGCTATTGAATTACAA

 15490 15500 15510 15520 15530 15540
 TACGTTGCAACCGATGAAAAATCATTACGGGAGATCTACCAGATTGGTTATGATGCTGC

 15550 15560 15570 15580 15590 15600
 AAAAGATCAACATGACAACCTTATGGCATTGAAAGAAAGTATCACCAACCAGCGAGGTTAA

 15610 15620 15630 15640 15650 15660
 AAAGAACGTCTTAGCAAATGGTTGGTATAACTGCTAGAACAGCGGCAAATAGCG

 15670 15680 15690 15700 15710 15720
 GCCCACACGGATTATACACTAGGATAATGGCGTTAATAGCCTCACTGTCGTTGTGG

 15730 15740 15750 15760 15770 15780
 TCTCTAATTTAGCTAAATCTTGTGTTACTGACTCCTATTAAATCATAAACGATTAT

 15790 15800 15810 15820 15830 15840
 CACGGTAAACATGACTCAAATAACCCGCTTCACGGCATGACACTCGAAAAAGTAAT

 15850 15860 15870 15880 15890 15900
 TAACAGTCTCGTTGAACAATATGGCTGGGATGGCTTGGATACTACATCAACATTGTTG

 15910 15920 15930 15940 15950 15960
 CTTTACTGAAAATCCAAGTGTAAAGTCTAGTCTAAATTTACGTAAAACCCCTTGGC

Fig. 5

AM

15970 15980 15990 16000 16010 16020
 ACGTGATAAAGTAGAAGCGCTATATATCAAAATGGTGAAGGCTAACTGTCTCCACG

 16030 16040 16050 16060 16070 16080
 CTAGCGAACCGCTGTTATAGTTAATATAAGTACTATAAGCAGGGCTCGTTAATTCAAGTA

 16090 16100 16110 16120 16130 16140
 TGTAATTAATCCTGAATACTCCGCTTATTCAACATGTACTCTCTAGATAACACTCTC

 16150 16160 16170 16180 16190 16200
 AACATTACACCTTCAACATCACAGCCTCCACATAACATCCGATGACATAGCCCTGTTATT

 16210 16220 16230 16240 16250 16260
 TTTCACATTTATCTATATGCTATATATTAGCCATTGATCAATTGAGTTAATTCTGC

 16270 16280 16290 16300 16310 16320
 AATGACAAAGATATACCATCATCCAGTACAAATTATTATGAAGATAACGACCATTCTGG

 16330 16340 16350 16360 16370 16380
 TGTTGTTTACCAACCTAACCTTTAAATACTTTGAACGTGCACGTGAGCATGTGATAAA

 16390 16400 16410 16420 16430 16440
 TAGTGACTTACTAGCAACATTGTGGAATGAACGCGTTAGGTTGCGGTGTATAAAGC

 16450 16460 16470 16480 16490 16500
 CAATATGACTTTCAAGGATGGGTCGAATTGCTGAAGTGTGTGATATTGCACTTCTTT

 16510 16520 16530 16540 16550 16560
 TGTCCTAGACGGTAAGTACAAACGATCTGGCGCCAAGAAGTATGGCGTCCGAATGCGAC

 16570 16580 16590 16600 16610 16620
 TAGGGCTGCCGTTATCGGTGATATTGAAATGGTGTGTTAGACAAACAAAACGTTTACA

 16630 16640 16650 16660 16670 16680
 GCCCATCCCTGATGATGTGTTAGCTGCAATGGTTAGTGAATAATGGTCATGCATAAAT

 16690 16700 16710 16720 16730 16740
 AGTTAACATGATTCTGGCCCGTCACGTTACAGATAAGAGGCATCCGATGCCTCCTTC

 16750 16760 16770 16780 16790 16800
 CTATTACCAATACTACTGCTTATCCCTTCTAACTATCTTAGCGTCCATAACACACTGA

 16810 16820 16830 16840 16850 16860
 GCATTATTCTATTAAATCAGTGATTGTGATTAAATTATCTTATATATGTAATTAAATG

 16870 16880 16890 16900 16910 16920
 TAATTTCATTTAGCTACATTAAGGCTTACGAATGTACGCTAAAATGAGATGT

 16930 16940 16950 16960 16970 16980
 CAGACTAATTTAGCTTATTAATCTGTTAGCCGTTATATTATAAAGATGGGATTAA

 16990 17000 17010 17020 17030 17040
 CTTAAATGCAATTAAATTATGGCGTAAATAGAGTGAAACATGGCTAAATATTCACTAAGTC

 17050 17060 17070 17080 17090 17100
 CTGAATTATATAAAGTTAATCTGTTATTAGCGTTACCTGGTCTTACAGTGAGG

Fig. 5

17110 17120 17130 17140 17150 17160
 TTTATGCCATTATTAGTGGGATTGAAGTGATTTAAAGCTATGTATATTATTGCAAAT

 17170 17180 17190 * 17200 17210 17220
 ATAAATTGTAACAATTAAGACTTGGACACTTGAGTTCAATTGAAATTGATTGGCATAA

 17230 17240 17250 17260 17270 17280
 AATTAAAACAGCTAAATCTACCTCAATCATTAGCAAATGTATGCAGGTAGATTTT

 17290 17300 17310 17320 17330 17340
 TCGCCATTAAGAGTACACTTGTACGCTAGGTTTTGTTAGTGTGCAAATGAACGTTT

 17350 17360 17370 17380 17390 17400
 GATGAGCATTGTTTAGAGCACAAAATAGATCCTTACAGGAGCAATAACGCAATGGCTA

 17410 17420 17430 17440 17450 17460
 AAAAGAACACCACATCGATTAAGCACGCCAAGGATGTTAAGTAGTAGTGATGATCAACAGT

 17470 17480 17490 17500 17510 17520
 TAAATTCTCGTTGCAAGAATGTCCGATTGCCATCATTGGTATGGCATCGGTTTTGCAG

 17530 17540 17550 17560 17570 17580
 ATGCTAAAAACTGGATCAATTCTGGATAACATCGTTGACTCTGTGGACGCTATTATG

 17590 17600 17610 17620 17630 17640
 ATGTGCCTAGCGATCGCTGGAACATTGACCGACCATTACTCGGCTGATAAAAAGCAGCTG

 17650 17660 17670 17680 17690 17700
 ACAAGACATACTGCAAACCGCGGTGGTTCATCAGAGCTGATTTGATCCGATGGAGT

 17710 17720 17730 17740 17750 17760
 TTGGTTTACCGCCAAATATCCTCGAGTTAAGTGTGACTGACATCGCTCAATTGTTGTCATTAATTG

 17770 17780 17790 17800 17810 17820
 TTGCTCGTGTATTAAAGTGTGACTGGCATTGGTAGTGATTATGACCATGATAAAATTG

 17830 17840 17850 17860 17870 17880
 GTATCACGCTGGGTGTCGGTGGTCAGAAACAAATTGCCATTAACGTCGCGCCTAC

 17890 17900 17910 17920 17930 17940
 AAGGCCCGGTATTAGAAAAAGTATTAAAGCCTCAGGCATTGATGAAGATGATCGCGCTA

 17950 17960 17970 17980 17990 18000
 TGATCATCGACAAATTAAAAAGCCTACATCGGCTGGGAAGAGAACTCATTCCCAGGCA

 18010 18020 18030 18040 18050 18060
 TGCTAGGTAACGTTATTGCTGGTCGTATGCCAATCGTTTGATTTGGTGGTACTAACT

 18070 18080 18090 18100 18110 18120
 GTGTGGTTGATGCCGATGCGCTGGCTCCCTGAGCTGTTAAATGGCGATCTCAGACT

 18130 18140 18150 18160 18170 18180
 TACTTGAATATCGTCAGAAGTCATGATATCGGGTGGTGTATGTTGTGATAACTGCCAT

 18190 18200 18210 18220 18230 18240
 TCATGTATATGTCATTCTGAAAACACCAGCAATTACCAATGATGATATCCGTCCGT

Fig. 5

18250 18260 18270 18280 18290 18300
 TTGATGACGATTCAAAAGGCATGCTGGTTGGTGAAGGTATTGGCATGATGGCGTTAAC

 18310 18320 18330 18340 18350 18360
 GTCTTGAAAGATGCTAACGTGACGGCGACAAAATTATTCTGTACTGAAAGGTATCGGTA

 18370 18380 18390 18400 18410 18420
 CATCTTCAGATGGTCGTTCAAATCTATTACGCTCCACGCCAGATGGCCAAGCAAAAG

 18430 18440 18450 18460 18470 18480
 CGCTAAAACGTGCTTATGAAGATGCCGGTTTGCCCTGAAACATGTGGTCTAATTGAAG

 18490 18500 18510 18520 18530 18540
 GCCATGGTACGGGTACCAAAGCGGGTGATGCCGCAGAATTGCTGGCTTGACCAAACACT

 18550 18560 18570 18580 18590 18600
 TTGGCGCCGCCAGTGATGAAAAGCAATATATCGCCTTAGGCTCAGTTAAATCGCAAATTG

 18610 18620 18630 18640 18650 18660
 GTCATACTAAATCTCGGGCTGGCTCTCGGGGTATGATTAAGGCGGCATTAGCGCTGCATC

 18670 18680 18690 18700 18710 18720
 ATAAAAATCTTACCTGCAACGATCCATATCGATAAACCAAGTGAAGCCTGGATATCAAAA

 18730 18740 18750 18760 18770 18780
 ACAGCCCGTTATACCTAAACAGCGAACCGTCCTGGATGCCACGTGAAGATGGTATT

 18790 18800 18810 18820 18830 18840
 CACGTCGTGCAGGTATCAGCTATTGGTTTGGCGGCACCAACTCCATATTATTTAG

 18850 18860 18870 18880 18890 18900
 AAGAGTATGCCAGGTACGATAGCGATATCGCTAAACTCAGTGAGCCAAACTGTGT

 18910 18920 18930 18940 18950 18960
 TGATCTCGGAAACGACCAACAAGGTATTGTTGCTGAGTTAAACTGGCGTACTAAC

 18970 18980 18990 19000 19010 19020
 TGGCTGTCGATGCTGATCATCAAGGGTTGTATTAATGAGTTAGTGACAAACGTGGCCAT

 19030 19040 19050 19060 19070 19080
 TAAAAACCCCATCCGTTAACCAAGCTCGTTAGGTTTGGCGCGTAATGCAAATGAAG

 19090 19100 19110 19120 19130 19140
 CGATCGCGATGATTGATAACGGCATTGAAACAAATTCAATGCGAACCGCAGATAAAATGACAT

 19150 19160 19170 19180 19190 19200
 GGTCACTACCGGGTTTACTATCGTCAAGCCCGTATTGATGCAACAGGTAAAGTGG

 19210 19220 19230 19240 19250 19260
 TTGCGCTATTCTCAGGGCAAGGTTCGCAATACGTGAACATGGGTGCGTGAATTAACCTGTA

 19270 19280 19290 19300 19310 19320
 ACTTCCCAAGCATGATGCACAGTGCTGCCGATGGATAAAGAGTTCAAGTGCCTGGTT

 19330 19340 19350 19360 19370 19380
 TAGGCCAGTTATCTGCAGTTACTTCCCTATCCCTGTTATACGGATGCCGAGCGTAAGC

Fig. 5

19390 19400 19410 19420 19430 19440
 TACAAGAAGAGCAATTACGTTAACGCAACATGCGCAACCAGCGATTGGTAGTTGAGTG

 19450 19460 19470 19480 19490 19500
 TTGGTCTGTTCAAAACGTTAACGCAAGCAGGTTAACGCTGATTGCTGCCGGTCATA

 19510 19520 19530 19540 19550 19560
 GTTTCGGTGAGTTAACCGCATTATGGCTGCCGATGTATTGAGCGAAAGCGATTACATGA

 19570 19580 19590 19600 19610 19620
 TGTTAGCGCGTAGTCGTGGTCAAGCAATGGCTGCCAGAGCAACAAGATTTGATGCAG

 19630 19640 19650 19660 19670 19680
 GTAAGATGGCCGCTGTTGGTATCCAAAGCAAGTCGCTGTGATCATTGATACCCCTG

 19690 19700 19710 19720 19730 19740
 ATGATGTCTCTATTGCTAACCTCAACTCGAATAACCAAGTTGTTATTGCTGGTACTACGG

 19750 19760 19770 19780 19790 19800
 AGCAGGTTGCTGTAGCGGTTACAACCTTAGGTAATGCTGGTTCAAAGTTGTGCCACTGC

 19810 19820 19830 19840 19850 19860
 CGGTATCTGCTGCCATACACCTTAGTTAGTCGTACGCGAAAAACCATTGCTAAAG

 19870 19880 19890 19900 19910 19920
 CGGTTGATAGCGCTAAATTAAAGCGCCAAGCATTCCAGTGTGTTGCTAACGGCACAGGCT

 19930 19940 19950 19960 19970 19980
 TGGTGCATTCAAGCAAACCGAATGACATTAAGAAAAACCTGAAAAACCACATGCTGGAAT

 19990 20000 20010 20020 20030 20040
 CTGTTCAATTCAATCAAGAAATTGACAAACATCTATGCTGATGGTGGCCCGTATTATCG

 20050 20060 20070 20080 20090 20100
 AATTTGGTCAAAGAATGTATTAACCAAATTGGTTGAAACATTCTCACTGAAAAATCTG

 20110 20120 20130 20140 20150 20160
 ATGTGACTGCTATCGCGGTTAATGCTAACCTAAACACCTGCGGACGTACAAATGCGCC

 20170 20180 20190 20200 20210 20220
 AAGCTGCGCTGCAAATGGCAGTGCTTGGTGTGCAATTAGACAATATTGACCCGTACGACG

 20230 20240 20250 20260 20270 20280
 CCGTTAACCGTCCACTTGGTGGCCGAAAGCATCACCAATGTTGATGAAGTTATCTGCAG

 20290 20300 20310 20320 20330 20340
 CGTCTTATGTTAGTCCGAAACGAAGAAAGCGTTGCTGATGCAATTGACTGATGGCTGGA

 20350 20360 20370 20380 20390 20400
 CTGTTAACGCAAGCGAAAGCTGTACCTGCTGTTGTCACAACCACAAAGTGATTGAAAAGA

 20410 20420 20430 20440 20450 20460
 TCGTTGAAAGTTGAAAAGATAGTTGAACGCATTGTCGAAGTAGAGCGTATTGTCGAAGTAG

 20470 20480 20490 20500 20510 20520
 AAAAACGTTACGTTAACGCTGACGGTTGCTTATATCGAAAATAATCAAGACGTTA

Fig. 5

20530 20540 20550 20560 20570 20580
 ACAGCGCTGTTAGCAACGTGACTAATAGCTCAGTGAECTAGCAGTGATGCTGACC

 20590 20600 20610 20620 20630 20640
 TTGTTGCCCTATTGAACGCAGTGTGGTCAATTGTTGCACACCAACAGCAATTATTAA

 20650 20660 20670 20680 20690 20700
 ATGTACATGAACAGTTATGCAAGGTCCACAAGACTACCGAAGAACAGTGCAGAACGTAC

 20710 20720 20730 20740 20750 20760
 TTGCTGCCAGACGAGCAATGAATTACCGGAAAGTTAGACCGTACATTGTCTATGTATA

 20770 20780 20790 20800 20810 20820
 ACGAGTTCCAATCAGAAACGCTACGTGTACATGAAACGTACCTGAACAATCAGACGAGCA

 20830 20840 20850 20860 20870 20880
 ACATGAACACCATGCTTACTGGTGCTGAAGCTGATGTGCTAGCAACCCCAATAACTCAGG

 20890 20900 20910 20920 20930 20940
 TAGTGAATACAGCCGTTGCCACTAGTCACAAGGTAGTTGCTCCAGTTATTGCTAATACAG

 20950 20960 20970 20980 20990 21000
 TGACGAATGTTGTATCTAGTGTAGTAATAACGCGGGTTGCAGTGCAAACGTGGCAT

 21010 21020 21030 21040 21050 21060
 TAGCGCCTACGCAAGAAATCGCTCCAACAGTCGCTACTACGCCAGCACCCGCATTGGTTG

 21070 21080 21090 21100 21110 21120
 CTATCGTGGCTGAACCTGTGATTGTTGCGCATGTTGCTACAGAAGTTGCACCAATTACAC

 21130 21140 21150 21160 21170 21180
 CATCAGTTACACCAGTTGTCGCAACTCAAGCGGCTATCGATGTAGCAACTATTAACAAAG

 21190 21200 21210 21220 21230 21240
 TAATGTTAGAAGTTGCTGATAAAACCGGTTATCCAACGGATATGCTGGAACGTGAGCA

 21250 21260 21270 21280 21290 21300
 TGGACATGGAAGCTGACTTAGGTATCGACTCAATCAAACGTGTTGAGATATTAGGCGCAG

 21310 21320 21330 21340 21350 21360
 TACAGGAATTGATCCCTGACTTACCTGAACCTAACCTGAAAGATCTGCTGAGCTACGCA

 21370 21380 21390 21400 21410 21420
 CGCTTGGTGAGATTGTCGATTACATGAATTCAAAAGCCCAGGCTGTAGCTCCTACAACAG

 21430 21440 21450 21460 21470 21480
 TACCTGTAACAAGTGCACCTGTTCGCCTGCATCTGCTGGTATTGATTAGCCCACATCC

 21490 21500 21510 21520 21530 21540
 AAAACGTAATGTTAGAAGTGGTTGCAGACAAAACCGGTTACCCAACAGACATGCTAGAAC

 21550 21560 21570 21580 21590 21600
 TGAGCATGGATATGGAAGCTGACTTAGGTATTGATTCAATCAAGCGTGTGGAAATCTTAG

 21610 21620 21630 21640 21650 21660
 GTGCAGTACAGGAGATCATAACTGATTACCTGAGCTAACCTGAAGATCTTGCTGAAT

Fig. 5

21670 21680 21690 21700 21710 21720
 TACGCACCCTAGGTGAAATCGTTAGTTACATGCAAAGCAAAGCGCCAGTCGCTGAAAGTG

 21730 21740 21750 * 21760 21770 21780
 CGCCAGTGGCGACGGCTCCTGTAGCAACAAGCTCAGCACCGTCTATCGATTGAAACCACA

 21790 21800 21810 21820 21830 21840
 TTCAAACAGTGTGATGGATGTAGTTGCAGATAAGACTGGTTATCCAATGACATGCTAG

 21850 21860 21870 21880 21890 21900
 AACTTGGCATGGACATGGAAGCTGATTAGGTATCGATTCAATCAAACGTGTGGAAATAT

 21910 21920 21930 21940 21950 21960
 TAGGCGCAGTGCAGGAGATCATCACTGATTACCTGAGCTAAACCCAGAAGACCTCGCTG

 21970 21980 21990 22000 22010 22020
 AATTACGCACGCTAGGTGAAATCGTTAGTTACATGCAAAGCAAAGCGCCAGTCGCTGAGA

 22030 22040 22050 22060 22070 22080
 GTGCGCCAGTAGCGACGGCTTCTGTAGCAACAAGCTCTGCACCGTCTATCGATTAAACC

 22090 22100 22110 22120 22130 22140
 ATATCCAAACAGTGTGATGGAAGTGGTTGCAGACAAAACCGGTTATCCAGTAGACATGT

 22150 22160 22170 22180 22190 22200
 TAGAACTTGCTATGGACATGGAAGCTGACCTAGGTATCGATTCAATCAAGCGTGTAGAAA

 22210 22220 22230 22240 22250 22260
 TTTTAGGTGCGGTACAGGAAATCATTACTGACTTACCTGAGCTTAACCCTGAAGATCTG

 22270 22280 22290 22300 22310 22320
 CTGAACTACGTACATTAGGTGAAATCGTTAGTTACATGCAAAGCAAAGCGCCCGTAGCTG

 22330 22340 22350 22360 22370 22380
 AAGCGCCTGCAGTACCTGTTGCAGTAGAAAGTGCACCTACTAGTGTAAACAGCTCAGCAC

 22390 22400 22410 22420 22430 22440
 CGTCTATCGATTAGACCACATCCAAAATGTAATGATGGATGTTGCTGATAAGACTG

 22450 22460 22470 22480 22490 22500
 GTTATCCTGCCAATATGCTTGAATTAGCAATGGACATGGAAGCCGACCTTGGTATTGATT

 22510 22520 22530 22540 22550 22560
 CAATCAAGCGTGTGAAATTCTAGGCGCGTACAGGAGATCATTACTGATTACCTGAAC

 22570 22580 22590 22600 22610 22620
 TAAACCCAGAAGACTTAGCTGAACCTACGTACGTTAGAAGAAATTGTAACCTACATGCAA

 22630 22640 22650 22660 22670 22680
 GCAAGGCGAGTGGTGTACTGTAATGTAGTGGCTAGCCCTGAAAATAATGCTGTATCAG

 22690 22700 22710 22720 22730 22740
 ATGCATTATGCAAAGCAATGTGGCGACTATCACAGCGGCCGAGAACATAAGGCGGAAT

 22750 22760 22770 22780 22790 22800
 TTAAACCGGCGCCGAGCGCAACCGTTGCTATCTCGTCTAAGCTCTACAGTAAAATAA

Fig. 5

22810 22820 22830 22840 22850 22860
 GCCAAGATTGTAAAGGTGCTAACGCCCTAACCGTAGCTGATGGCACTGATAATGCTGTGT

 22870 22880 22890 22900 22910 22920
 TACTTGCAGACCACCTATTGCAAACCTGGCTGGAAATGTAACTGCAATTGCAACCAACTTGGG

 22930 22940 22950 22960 22970 22980
 TAGCTGTAACAACGACGAAAGCATTAAAGTCAGTGAACCTGGTGACTTTAAATGGCG

 22990 23000 23010 23020 23030 23040
 TTGATGAAACTGAAATCAACAAACATTATTACTGCTAACGCACAATTGGATGCAGTTATCT

 23050 23060 23070 23080 23090 23100
 ATCTGCACGCAAGTAGCAGAAATTAAATGCTATCGAATACCCACAAGCATCTAACGCAAGGCC

 23110 23120 23130 23140 23150 23160
 TGATGTTAGCCTTCTTATTAGCGAAATTGAGTAAAGTAACCTAACGGCGCTAAAGTGCCTG

 23170 23180 23190 23200 23210 23220
 GCGCCTTATGATTGTTACTCAGCAGGGTGGTCATTAGGTTTGATGATATCGATTCTG

 23230 23240 23250 23260 23270 23280
 CTACAAGTCATGATGTGAAAACAGACCTAGTACAAAGCGGCTTAAACGGTTAGTTAAGA

 23290 23300 23310 23320 23330 23340
 CACTGTCTCACGAGTGGGATAACGTATTCTGTCGTGCGGTTGATATTGCTTCGTCTAA

 23350 23360 23370 23380 23390 23400
 CGGCTGAACAAGTTGCAAGCCTGTTAGTGAACACTTGATGCTAACACTGTATTAA

 23410 23420 23430 23440 23450 23460
 CAGAAGTGGTTATCAACAAGCTGGTAAAGGCCTTGAACGTATCACGTTACTGGTG

 23470 23480 23490 23500 23510 23520
 CTACTGACAGCTATGCATTAACAGCTGGCAATAACATCGATGCTAACTCGGTATTTAG

 23530 23540 23550 23560 23570 23580
 TGAGTGGTGGCGCAAAGGTGTAAGTGCACATTGTTGCTCGTATAGCTAAAGAATATC

 23590 23600 23610 23620 23630 23640
 AGTCTAAGTTCATCTTATTGGGACGTTCAACGTTCTCAAGTGACGAACCGAGCTGGCAA

 23650 23660 23670 23680 23690 23700
 GTGGTATTACTGATGAAGCGGCGTTAAAGAAAGCAGCGATGCAGTCTTGATTACAGCAG

 23710 23720 23730 23740 23750 23760
 GTGATAAACCAACACCCGTTAACGATCGTACAGCTAACCAACCAATCCAAGCTAACCGTG

 23770 23780 23790 23800 23810 23820
 AAATTGCGCAAACCTTGTCTGCAATTACCGCTGCTGGTGGCCAAGCTGAATATGTTCTG

 23830 23840 23850 23860 23870 23880
 CAGATGTAACTAATGCAGCAAGCGTACAAATGGCAGTCGCTCCAGCTATCGCTAACGTC

 23890 23900 23910 23920 23930 23940
 GTGCAATCACTGGCATTCATGGCGCGGGTGTAGCTGACCAATTGAGCAAA

Fig. 5

23950 23960 23970 23980 23990 24000
 AAACACTGAGTGATTTGAGTCTGTTACAGCACTAAAATTGACGGTTGTTATCGCTAC

 24010 24020 24030 24040 24050 24060
 TATCAGTCACTGAAGCAACATCAAGCAATTGGTATTGTTCTCGTCAGCGGCTGGTT

 24070 24080 24090 24100 24110 24120
 TCTACGGTAACCCCGGCCAGTCTGATTACTCGATTGCCAATGAGATCTAAATAAAACCG

 24130 24140 24150 24160 24170 24180
 CATAACCGCTTTAAATCATTGCACCCACAAGCTCAAGTATTGAGCTTAACGGGTCCTT

 24190 24200 24210 24220 24230 24240
 GGGACGGTGGCATGGTAACGCCTGAGCTTAAACGTATGTTGACCAACGTGGTGTAA

 24250 24260 24270 24280 24290 24300
 TTATTCCACTTGATGCAGGTGCACAGTTATTGCTGAATGAACTAGCCGCTAATGATAACC

 24310 24320 24330 24340 24350 24360
 GTTGTCCACAAATCCTCGTGGGTAAATGACTTATCTAAAGATGCTAGCTCTGATCAAAAGT

 24370 24380 24390 24400 24410 24420
 CTGATGAAAAGAGTACTGCTGTAAAAAGCCACAAGTTAGTCGTTATCAGATGCTTAG

 24430 24440 24450 24460 24470 24480
 TAACTAAAAGTATCAAAGCGACTAACAGTAGCTCTTATCAAACAAGACTAGTGCTTAT

 24490 24500 24510 24520 24530 24540
 CAGACAGTAGTGCTTTCAAGGTTAACGAAAACCACTTTAGCTGACCACATGATCAAAG

 24550 24560 24570 24580 24590 24600
 GCAATCAGGTATTACCAACGGTATGCGCGATTGCTGGATGAGTGATGCAGCAAAAGCGA

 24610 24620 24630 24640 24650 24660
 CTTATAGTAACCGAGACTGTGCATTGAAGTATGTCGGTTCGAAGACTATAAATTGTTA

 24670 24680 24690 24700 24710 24720
 AAGGTGTGGTTTGATGGCAATGAGGCAGGCGATTACCAAATCCAATTGTCGCCTGTGA

 24730 24740 24750 24760 24770 24780
 CAAGGGCGTCAGAACAGGATTCTGAAGTCCGTATTGCCGAAAGATCTTAGCCTGAAAA

 24790 24800 24810 24820 24830 24840
 GTGACGGTAAACCTGTGTTCAATTATGCAGCGACAATTGTTAGCAACTCAGCCACTTA

 24850 24860 24870 24880 24890 24900
 ATGCTGTGAAGGTAGAACTTCCGACATTGACAGAAAGTGTGATAGCAACAATAAAGTAA

 24910 24920 24930 24940 24950 24960
 CTGATGAAGCACAAGCGTTACAGCAATGGCACCTGTTCCACGGTAAAGTCTGCAGG

 24970 24980 24990 25000 25010 25020
 GCATTAAGCAGATATTAAGTTGTGACGACAAGGGCCTGCTATTGGCTGTAGATAACCG

 25030 25040 25050 25060 25070 25080
 ATGTTGCAACAGCTAACGAGGGATCCTCCCGTTAGCTGACAACAATATCTTGCCAATG

Fig. 5

25090 25100 25110 25120 25130 25140
 ATTTGGTTATCAGGCTATGTTGGCTGGGTGCGCAAACAATTGGTTAGGTAGCTTAC

 25150 25160 25170 25180 25190 25200
 CTTGGTGACAACGGCTTGGACTGTGTATCGTGAAGTGGTTGTAGATGAAGTATTTCATC

 25210 25220 25230 25240 25250 25260
 TGCAACTTAATGTTGGTGGCATGATCTATTGGGTCACGCCAGTAAAGGCCGTTGTG

 25270 25280 25290 25300 25310 25320
 ATATTCAATTGATTGCTGCTGATATGCAATTACTGCCGAAGTGAAATCAGCGCAAGTCA

 25330 25340 25350 25360 25370 25380
 GTGTCAGTGACATTGAAACGATATGTCATGATCGAGTAAATAAACGATAGGCGTCAT

 25390 25400 25410 25420 25430 25440
 GGTGAGCATGGCGTCTGCTTCTCATTTTAACATTAACAATATTAATAGCTAACGC

 25450 25460 25470 25480 25490 25500
 GGTTGCTTAAACCAAGTAAACAAGTGCTTTAGCTATTACTATTCAAACAGGATATTA

 25510 25520 25530 25540 25550 25560
 AAGAGAATATGACGGAATTAGCTGTTATTGGTATGGATGCTAAATTAGCGGACAAGACA

 25570 25580 25590 25600 25610 25620
 ATATTGACCGTGTGGAACCGCGCTTCTATGAAGGTGCTTATGTAGGTAATGTTAGCCCG

 25630 25640 25650 25660 25670 25680
 TTAGTACCGAATCTAATGTTATTAGCAATGGCGAAGAACAAAGTTATTACTGCCATGACAG

 25690 25700 25710 25720 25730 25740
 TTCTTAACCTCTGTCAGTCTACTAGCGCAAACGAATCAGTTAAATATAGCTGATATCGCG

 25750 25760 25770 25780 25790 25800
 TGTTGCTGATTGCTGATGTAAGGTGCTGATGATCAGCTTGTAGTCCAAATTGCATCAG

 25810 25820 25830 25840 25850 25860
 CAATTGAAAAACAGTGTGCGAGTTGTGTTATTGCTGATTAGGCCAAGCATTAAATC

 25870 25880 25890 25900 25910 25920
 AAGTAGCTGATTAGTTAAACCAAGACTGTCCTGGCTGTAATTGGCATGAATAACT

 25930 25940 25950 25960 25970 25980
 CGGTTAATTATCTCGTCATGATCTGAATCTGTAAGTCAACAAATCAGCTTGATGAAA

 25990 26000 26010 26020 26030 26040
 CCTTCAATGGTTATAACAATGAGCTGGGTTCGCGAGTTACTTATCGCTCAACTGCGT

 26050 26060 26070 26080 26090 26100
 TTGCCAATGCTAAGCAATGTTATATACGCCAACATTAAGGGCTTCGCTCAATCGGGCG

 26110 26120 26130 26140 26150 26160
 TAAATGCTCAATTAAACGTTGGAAACATTAGCGATACTGCAAAGACCGCATTGCAGCAAG

 26170 26180 26190 26200 26210 26220
 CTAGCATAACTGCAGAGCAGGTTGGTTGTTAGAAGTGTCAAGCAGTCGCTGATTGGCAA

Fig. 5

26230 26240 26250 26260 26270 26280
 TCGCATTGCTGAAAGCCAAGGTTAACATGCTGTTATCATACGCAAACCTTGCATA

 26290 26300 26310 26320 26330 26340
 CTGCATTAAGCAGTGCCGTAGTGTGACTGGTGAAGGCGGGTGTTCACAGGTCGCAG

 26350 26360 26370 26380 26390 26400
 GTTTATTGAAATGTGTAATTGGTTACATCAACGTTATTCGGCGATTAAAGATTGGC

 26410 26420 26430 26440 26450 26460
 AACAAACCGAGTGACAATCAAATGTCACGGTGGCGGAATTCAACCATTCTATATGCCGTAG

 26470 26480 26490 26500 26510 26520
 ATGCTCGACCTGGTCCCACATGCTGATGGCTCTGCACACATTGCCGCTTATAGTTGTG

 26530 26540 26550 26560 26570 26580
 TGACTGCTGACAGCTATTGTCATATTCTTTACAAGAAAACGTCTTACAAGAAACTTGT

 26590 26600 26610 26620 26630 26640
 TGAAAGAAACAGTCTGCAAGATAATGACTTAACGTGAAAGCAAGCTTCAGACTCTGAAC

 26650 26660 26670 26680 26690 26700
 AAAACAATCCAGTAGCTGATCTGCGCACTAATGGTTACTTGCATCGAGCGAGTTAGCAT

 26710 26720 26730 26740 26750 26760
 TAATCATAGTACAAGGTAATGACGAAGCACAATTACGCTGTGAATTAGAAACTATTACAG

 26770 26780 26790 26800 26810 26820
 GGCAGTTAAGTACTACTGGCATAAGTACTATCAGTATTAAACAGATCGCAGCAGACTGTT

 26830 26840 26850 26860 26870 26880
 ATGCCCGTAATGATACTAACAAAGCCTATAGCGCAGTGCTTATGCCGAGACTGCTGAAG

 26890 26900 26910 26920 26930 26940
 AGTTAAGCAAAGAAATAACCTGGCGTTGCTGGTATCGCTAGCGTGTAAATGAAGATG

 26950 26960 26970 26980 26990 27000
 CTAAAGAATGGAAAACCCCGAAGGGCAGTTATTTACCGCGCAGCCTGCAAATAAACAGG

 27010 27020 27030 27040 27050 27060
 CTGCTAACAGCACACAGAACATGGTGTACCTCATGTACCCAGGTATTGGTGCTACATATG

 27070 27080 27090 27100 27110 27120
 TTGGTTAGGGCGTGATCTATTCATCTATTCCCACAGATTATCAGCCTGTAGCGGCTT

 27130 27140 27150 27160 27170 27180
 TAGCCGATGACATTGGCGAAAGTCTAAAGATACTTACTTAATCCACGCAGTATTAGTC

 27190 27200 27210 27220 27230 27240
 GTCATAGCTTAAAGAACTCAAGCAGTTGGATCTGGACCTGCGCGGTAACTTAGCCAATA

 27250 27260 27270 27280 27290 27300
 TCGCTGAAGCCGGTGTGGTTTGCTTGTGTTACCAAGGTATTGAAGAAGTCTTTG

 27310 27320 27330 27340 27350 27360
 CCGTTAAAGCTGACTTGTACAGGTTATAGCATGGGTGAAGTAAGCATGTATGCAGCAC

Fig. 5

27370 27380 27390 27400 27410 27420
 TAGGCTGCTGGCAGCAACCGGGATTGATGAGTGCTCGCCTTGACAAATCGAACATACCTTA

 27430 27440 27450 27460 27470 27480
 ATCATCAACTTTGCCGGAGTTAAGAACACTACGTCAAGCATTGGGCATGGATGATGTAG

 27490 27500 27510 27520 27530 27540
 CTAACGGTACGTTCGAGCAGATCTGGGAAACCTATACCATTAAGGCAACGATTGAACAGG

 27550 27560 27570 27580 27590 27600
 TCGAAATTGCCTCTGCAGATGAAGATCGTGTATTGCACCATTATCAATACACCTGATA

 27610 27620 27630 27640 27650 27660
 GCTTGTGTTAGCCGGTTATCCAGAAGCCTGTCAGCGAGTCATTAAGAATTAGGTGTGC

 27670 27680 27690 27700 27710 27720
 GTGCAATGGCATTGAATATGGCGAACGCAATTACAGCGGCCAGCTTATGCCGAATACG

 27730 27740 27750 27760 27770 27780
 ATCATATGGTTGAGCTATACCATATGGATGTTACTCCACGTATTAATACCAAGATGTATT

 27790 27800 27810 27820 27830 27840
 CAAGCTCATGTTATTACCGATTCCACAACGCAGCAAAGCGATTCCCACAGTATTGCTA

 27850 27860 27870 27880 27890 27900
 AATGTTGTGATGTGGATTCCACGTTGGTTAACCTTACATGACAAAGGTG

 27910 27920 27930 27940 27950 27960
 CGCGGGTATTCAATTGAAATGGGTCCAGGTGTTCTGTTATGTAGCTGGTAGATAAGATCT

 27970 27980 27990 28000 28010 28020
 TAGTTAATGGCGATGGCGATAATAAAAGCAAAGCCAACATGTATCTGTCCTGTGAATG

 28030 28040 28050 28060 28070 28080
 CCAAAGGCACCAAGTGAACTTACTTATATTGCGATTGCTAACGTTAATTAGTCATG

 28090 28100 28110 28120 28130 28140
 GCGTGAATTGAAATTAGATAGCTTGTAAACGGGTCACCTGGTTAACAGCAGGCCATA

 28150 28160 28170 28180 28190 28200
 TAGCAAACACGAACAAATAGTCAACATCGATATCTAGCGCTGGTGAGTTACCTCATTA

 28210 28220 28230 28240 28250 28260
 GTTGAATATGGATTTAAAGAGAGTAATTATGGAAAATATTGCAAGTAGTAGGTATTGCTA

 28270 28280 28290 28300 28310 28320
 ATTTGTTCCGGGCTACAAGCACCGGATCAATTGGCAGCAATTGCTGAACAAACAG

 28330 28340 28350 28360 28370 28380
 ATTGCCGAGTAAGGCAGCGCTGTTCAAATGGCGTTGATCCTGCTAACATACCGCCA

 28390 28400 28410 28420 28430 28440
 ACAAAAGGTGACACAGATAAAATTACTGTGTGCACGGCGGTTACATCAGTGATTCAATT

 28450 28460 28470 28480 28490 28500
 TTGATGCTTCAGGTTATCAACTCGATAATGATTATTAGCCGGTTAGATGACCTTAATC

Fig. 5

28510 28520 28530 28540 28550 28560
 AATGGGGGCTTTATGTTACGAAACAAGCCCTTACCGATGCGGGTATTGGGGCAGTACTG

 28570 28580 28590 28600 28610 28620
 CACTAGAAAACGTGGTGTGATTTAGGTAAATTGTCATTCCAACTAAATCATCTAACATC

 28630 28640 28650 28660 28670 28680
 AGCTGTTATGCCTTGTATCATCAAGTTGGTATAATGCCTTAAAGGCGGTATTACATC

 28690 28700 28710 28720 28730 28740
 CTGATTTCAATTAACGCATTACACAGCACCGAAAAAAACACATGCTGACAATGCATTAG

 28750 28760 28770 28780 28790 28800
 TAGCAGGTTATCCAGCTGCATTGATCGCGCAAGCGGCCGGTCTGGTGGTTCACATTTG

 28810 28820 28830 28840 28850 28860
 CACTGGATGCGGCTTGTGCTTCATCTTGTATAGCGTTAAGTTAGCGTGTGATTACCTGC

 28870 28880 28890 28900 28910 28920
 ATACGGGTAAAGCCAACATGATGCTTGCTGGTGGTATCTGCAGCAGATCCTATGTCG

 28930 28940 28950 28960 28970 28980
 TAAATATGGGTTCTCGATATTCCAAGCTTACCCAGCTAACATGTACATGCCCGTTG

 28990 29000 29010 29020 29030 29040
 ACCAAAATTCAACAAGGTCTATTGCCGGTGAAGGCCGGCATGATGGTATTGAAACGTC

 29050 29060 29070 29080 29090 29100
 AAAGTGATGCAGTACGTGATGGTATCATTTACGCCATTATTAAAGGCCGGCGCATTAT

 29110 29120 29130 29140 29150 29160
 CGAATGACGGTAAAGGCAGTTGTATTAAGCCCACACCAAGGGCCAAGTATTAGTAT

 29170 29180 29190 29200 29210 29220
 ATGAACGTGCTTATGCCGATGCAGATGTTGACCCGAGTACAGTTGACTATATTGAATGTC

 29230 29240 29250 29260 29270 29280
 ATGCAACGGGCACACCTAACGGGTACAATGTTGAATTGCGTTCGATGGAAACCTTTCA

 29290 29300 29310 29320 29330 29340
 GTCGCGTAAATAACAAACCATTACTGGGCTGGTTAAATCTAACCTGGTCATTGTTAA

 29350 29360 29370 29380 29390 29400
 CTGCCGCTGGTATGCCCTGGCATGACCAAAGCTATGTTAGCGCTAGGTAAAGGTCTTATTCA

 29410 29420 29430 29440 29450 29460
 CTGCAACGATTAACCTAAAGCAACCACTGCAATCTAAACCGTTACTTACTGGCGAGC

 29470 29480 29490 29500 29510 29520
 AAATGCCAACGACGACTGTGCTTGGCCAACAACTCCGGGTGCCAAGGCAGATAAACCGC

 29530 29540 29550 29560 29570 29580
 GTACCGCAGGTGTGAGCGTATTGGTTGGTGGCAGCAACGCCATTGGTATTACAAC

 29590 29600 29610 29620 29630 29640
 AGCCAACGCAAACACTCGAGACTAATTAGTGTGCTAAACCACGTGAGCCTTGGCTA

Fig. 5

29650 29660 29670 29680 29690 29700
 TTATTGGTATGGACAGCCATTTGGTAGTGCCAGTAATTAGCGCAGTTCAAAACCTTAT

 29710 29720 29730 29740 29750 29760
 TAAATAATAATCAAAATACCTCCGTGAATTACCAAGAACACGCTGGAAAGGCATGGAAA

 29770 29780 29790 29800 29810 29820
 GTAACGCTAACGTCATGCAGTCGTTACAATTACGCAAAGCGCCTAAAGGCAGTTACGTG

 29830 29840 29850 29860 29870 29880
 AACAGCTAGATATTGATTCTTGCCTTAAAGTACCGCCTAATGAAAAAGATTGCTTGA

 29890 29900 29910 29920 29930 29940
 TCCCGCAACAGTTAATGATGATGCAAGTGGCAGACAATGCTGCGAAAGACGGAGGTCTAG

 29950 29960 29970 29980 29990 30000
 TTGAAGGTCGTAATGTTGCGGTATTAGTAGCGATGGGCATGGAACATTACATCAGT

 30010 30020 30030 30040 30050 30060
 ATCGTGGTCGCCTTAATCTAACCAACCCAAATTGAAGACAGCTTATTACAGCAAGGTATTA

 30070 30080 30090 30100 30110 30120
 ACCTGACTGTTGAGCAACGTGAAGAACTGACCAATATTGCTAAAGACGGTGTGCCTCGG

 30130 30140 30150 30160 30170 30180
 CTGCACAGCTAAATCAGTATACGAGTTCTGGTAATATTATGGCGTCACGTATTCGG

 30190 30200 30210 30220 30230 30240
 CGTTATGGGATTTCTGGCCTGCTATTACCGTATCGGCTGAAGAAAACCTCTGTTATC

 30250 30260 30270 30280 30290 30300
 GTTGTGTTGAATTAGCTGAAAATCTATTCAAACCCAGTGATGTTGAAGCCGTTATTATTG

 30310 30320 30330 30340 30350 30360
 CTGCTGTTGATTTGCTGGTTCAATTGAAAACATTACTTACGTCAGCACTACGGTCAG

 30370 30380 30390 30400 30410 30420
 TTAATGAAAAGGGATCTGTAAGTGAATGTGGTCCGGTTAATGAAAGCAGTTCACTAACCA

 30430 30440 30450 30460 30470 30480
 ACAATATTCTTGATCAGCAACAAATGGCTGGTGGGTGAAGGCGCAGCGGCTATTGTCGTTA

 30490 30500 30510 30520 30530 30540
 AACCGTCATCGCAAGTCACTGCTGAGCAAGTTATGCGCGTATTGATGCGGTGAGTTTG

 30550 30560 30570 30580 30590 30600
 CCCCTGGTAGCAATGCGAAAGCAATTACGATTGCAGCGGATAAAGCATTAAACACTTGCTG

 30610 30620 30630 30640 30650 30660
 GTATCAGTGCTGCTGATGTTAGCTAGTGTTGAAGCACATGCAAGTGGTTAGTGCCGAAA

 30670 30680 30690 30700 30710 30720
 ATAATGCTGAAAAACCGCGTTACCGACTTATACCCAAGCGCAAGTATCAGTTCGGTGA

 30730 30740 30750 30760 30770 30780
 AAGCCAATATTGGTCATACGTTAATGCCTCGGGTATGGCGAGTATTAAAACGGCGC

Fig. 5

30790 30800 30810 30820 30830 30840
 TGCTGTTAGATCAGAATACTGAGTCAGAGATCAGAAAAGCAAACATATTGCTATTAACGGTC

 30850 30860 30870 30880 30890 30900
 TAGGTCGTATAACAGCTGCGCCATCTTATCTGAGTCAGCGAAGCGCATCAAG

 30910 30920 30930 30940 30950 30960
 TTGCACCAAGCGCCTGTATCTGGTATGGCCAAGCAACGCCACAGTTAGTTAAAACCATCA

 30970 30980 30990 31000 31010 31020
 AACTCGGTGGTCAGTTAATTAGCAACGCGATTGTTAACAGTGCGAGTTCATCTTACACG

 31030 31040 31050 31060 31070 31080
 CTATTAAAGCGCAGTTGCCGGTAAGCACTTAAACAAAGTTAACCGCCAGTGATGATGG

 31090 31100 31110 31120 31130 31140
 ATAACCTGAAGCCCCAAGGTATTAGCGCTATGCAACCAATGAGTATGTGGTACTGGAG

 31150 31160 31170 31180 31190 31200
 CTGCTAACACTCAAGCTTCTAACATTCAAGCATCTCATGTTCAAGCGTCAAGTCATGCCAC

 31210 31220 31230 31240 31250 31260
 AAGAGATAGCACCAAACCAAGTTCAAAATATGCAAGCTACAGCAGCCGCTGTAAGTTCAC

 31270 31280 31290 31300 31310 31320
 CCCTTTCTCAACATCAACACACAGCGCAGCCCGTAGCGGCACCGAGCGTTGGAGTGA

 31330 31340 31350 31360 31370 31380
 CTGTGAAACATAAAGCAAGTAACCAAATTCATCAGCAAGCGTCTACGCATAAAAGCATT

 31390 31400 31410 31420 31430 31440
 TAGAAAAGTCGTTAGCTGCACAGAAAAACCTATCGCAACTTGTGAATTGCAAACCAAGC

 31450 31460 31470 31480 31490 31500
 TGTCAATCCAAACTGGTAGTGACAATACTAACAATACTGCGTCAACAAGCAATACAG

 31510 31520 31530 31540 31550 31560
 TGCTAACAAATCCTGTATCAGCAACGCCATTAACACTTGTGCTAATGCCCTGTAGTAG

 31570 31580 31590 31600 31610 31620
 CGACAAACCTAACCAAGTACAGAAGCAAAAGCGCAAGCAGCTGCTACACAAGCTGGTTTC

 31630 31640 31650 31660 31670 31680
 AGATAAAAGGACCTGTTGGTTACAACATCCACCGCTGCAGTTAATTGAACGTTATAATA

 31690 31700 31710 31720 31730 31740
 AACCAAGAAAACGTGATTACGATCAAGCTGATTGGTTGAATTGCTGAAGGTGATATTG

 31750 31760 31770 31780 31790 31800
 GTAAGGTATGGTGTGAATACAATATTATTGATGGCTATTGCGCGTGTACGTCTGC

 31810 31820 31830 31840 31850 31860
 CAACCTCAGATTACTGTTAGTAACACGTGTTACTGAACCTGATGCCAAGGTGCATGAAT

 31870 31880 31890 31900 31910 31920
 ACAAGAAATCATACATGTGTACTGAATATGATGTGCCTGTTGATGCACCGTTCTTAATTG

Fig. 5

31930 31940 31950 31960 31970 31980
 ATGGTCAGATCCCTGGTCTGTTGCCGTCGAATCAGGCCAGTGTGATTTGATGTTGATT

 31990 32000 32010 32020 32030 32040
 CATATATCGGTATTGATTCCAAGCGAAAGGCGAACGTGTTACCGTTACTTGATTGTG

 32050 32060 32070 32080 32090 32100
 AATTAACCTTCCTTGAAGAGATGGCTTGGTGGCGATACTTACGTTACGAGATCCACA

 32110 32120 32130 32140 32150 32160
 TTGATTCGTATGCACGTAACGGCGAGCAATTATTATTCTTCCATTACGATTGTTACG

 32170 32180 32190 32200 32210 32220
 TAGGGGATAAGAAGGTACTTATCATGCGTAATGGTTGTGCTGGTTCTTACTGACGAAG

 32230 32240 32250 32260 32270 32280
 AACTTTCTGATGGTAAAGGCATTACATAACGACAAAGACAAAGCTGAGTTAGCAATG

 32290 32300 32310 32320 32330 32340
 CTGTTAAATCATCATCAGCGCTTATTACAACATAACCGTGGTCAATACGATTATAACG

 32350 32360 32370 32380 32390 32400
 ACATGATGAAGTTGGTTAATGGTGATGTTGCCAGTTGGTCCGCAATATGATCAAG

 32410 32420 32430 32440 32450 32460
 GTGGCCGTAATCCATCATTGAAATTCTCGTCTGAGAAGTTCTGATGATTGAACGTATTA

 32470 32480 32490 32500 32510 32520
 CCAAGATAGACCCAACCGGTGGTCATTGGGGACTAGGCCTGTTAGAAGGTAGAAAGATT

 32530 32540 32550 32560 32570 32580
 TAGACCTGAGCATTGGTATTCCCTTGTCACTTAAAGGTGATCAAGTAATGGCTGGTT

 32590 32600 32610 32620 32630 32640
 CGTTGATGTCGGAAGGTTGTGCCAAATGGCGATGTTCTCATGCTCTTGGTATGC

 32650 32660 32670 32680 32690 32700
 ATACCAATGTGAACAAACGCTCGTTCCAACCACTACCAAGGTGAATCACAAACGGTACGTT

 32710 32720 32730 32740 32750 32760
 GTCGTGGCAAGTACTGCCACAGCGCAATACCTAACCGTATGGAAGTTACTGCGA

 32770 32780 32790 32800 32810 32820
 TGGGTATGCATCCACAGCCATTGAAAGCTAATATTGATATTTGCTTGACGGTAAAG

 32830 32840 32850 32860 32870 32880
 TGGTTGTTGATTTCAAAACTTGAGCGTGATGATCAGCGAACAGATGAGCATTGAGATT

 32890 32900 32910 32920 32930 32940
 ACCCTGTAACACTGCCGAGTAATGTGGCGCTTAAAGCGATTACTGCACCTGTTGCGTCAG

 32950 32960 32970 32980 32990 33000
 TAGCACCAGCATCTCACCCGCTAACAGCGCGGATCTAGACGAACGTGGTGTGAACCGT

 33010 33020 33030 33040 33050 33060
 TTAAGTTCCCTGAACGTCCGTTAATGCGTGTGAGTCAGACTTGTCTGCACCGAAAAGCA

Fig. 5

33070 33080 33090 33100 33110 33120
 AAGGTGTGACACCGATTAAGCATTGAGCGCCTGCTGTTGGTCATCAGAGTGC

 33130 33140 33150 33160 33170 33180
 CTAACCAAGCACCCTTACACCTTGGCATATGTTGAGTTGCGACGGTAATATTCTA

 33190 33200 33210 33220 33230 33240
 ACTGTTTCGGTCCTGATTTGATGTTATGAAGGTCGTATTCCACCTCGTACACCTTGTG

 33250 33260 33270 33280 33290 33300
 GCGATTACAAGTTGTTACTCAGGTTGAGAAGTGCAGGGCGAACGTCTTGATCTTAAA

 33310 33320 33330 33340 33350 33360
 ATCCATCAAGCTGTGATGCTGAATACTATGTACCGGAAGACGCTTGGTACTTTACTAAA

 33370 33380 33390 33400 33410 33420
 ACAGCCATGAAAATGGATGCCTTATTCAATTACATGGAAATTGCATTGCAACCAAATG

 33430 33440 33450 33460 33470 33480
 GCTTTATTCTGGTTACATGGCACGACGCTTAAATACCCCTGAAAAAGATCTGTTCTTCC

 33490 33500 33510 33520 33530 33540
 GTAACCTTGATGGTAGCGGCACGTTATTAAAGCAGATTGATTTACGCGGCAAGACCATTG

 33550 33560 33570 33580 33590 33600
 TGAATAAAATCAGTCTGGTTAGTACGGCTATTGCTGGTGGCGCGATTATTCAAAGTTCA

 33610 33620 33630 33640 33650 33660
 CGTTTGATATGTCTGTAGATGGCGAGCTATTTATACTGGTAAAGCTGTATTGGTTACT

 33670 33680 33690 33700 33710 33720
 TTAGTGGTGAATCACTGACTAACCAACTGGCATTGATAACGGTAAAACGACTAATGCGT

 33730 33740 33750 33760 33770 33780
 GGTTTGGTATAACAATACCCCCGCAGCGAATTGATGTGTTGATTAACGTTAATCAGT

 33790 33800 33810 33820 33830 33840
 CATTGGCTCTGTATAAAGCGCCTGTTGAGAAACCGCATTATAAATTGGCTGGTGGTCAGA

 33850 33860 33870 33880 33890 33900
 TGAACCTTATCGATACAGTGTCACTGGTTGAAGGCGGTGGTAAAGCGGGCGTGGCTTATG

 33910 33920 33930 33940 33950 33960
 TTTATGGCGAACGTACGATTGATGCTGATGATTGGTTCTTCCGTTATCACTCCACCAAG

 33970 33980 33990 34000 34010 34020
 ATCCGGTGATGCCAGGTTCAATTAGGTGTTGAAGCTATTATTGAGTTGATGCAAGACCTATG

 34030 34040 34050 34060 34070 34080
 CGCTTAAAAATGATTGGTGGCAAGTTGCTAACCCACGTTTCAATTGCGCCGATGACGC

 34090 34100 34110 34120 34130 34140
 AAGTTGATTGGAAATACCGTGGCAAATTACGCCGCTGAATAAACAGATGTCACTGGACG

 34150 34160 34170 34180 34190 34200
 TGCAATCACTGAGATCGTGAATGACGCTGGTGAAGTGCAGATCGTGGTATGCGAATC

Fig. 5

34210 34220 34230 34240 34250 34260
 TGTCTAAAGATGGTCTCGTATTATGAAGTTAAAACATCGTTAAGTATTGTTGAAG

 34270 34280 34290 34300 34310 34320
 CGTAAAGGGTCAAGTGTAAACGTGCTTAAGCGCCGCATTGGTTAAAGACGCTTGCACGCC

 34330 34340 34350 34360 34370 34380
 GTGAATCCGTCCATGGAGGCTTGGGTTGGCATCCATGCCAACACAGCAAGCTTACTTT

 34390 34400 34410 34420 34430 34440
 AATCAATAACGGCTTGGTGTCCATTAGACGCCCTCGAACTTAGTTAATAGACAAAATA

 34450 34460 34470 34480 34490 34500
 ATTTAGCTGTGGAATGAATATAGTAAGTAATCATTGGCAGCTACAAAAAAGGAATTAAG

 34510 34520 34530 34540 34550 34560
 AATGTCGAGTTAGGTTAACAAATAACAGCAATTAACTGGGTTGGAAAGTAGATCC

 34570 34580 34590 34600 34610 34620
 AGCGTCAGTCATACACAAGATGCAGAAATTAAAGCAGCTTAACTGGATCTAACTAAACC

 34630 34640 34650 34660 34670 34680
 TCTCTATGTGGCGAATAATTCAAGCGTAACGGTATAGCTAATCATACTACGTCAAGTAGCAGG

 34690 34700 34710 34720 34730 34740
 TGCGATCAGCAATAACATCGATGTTGATGTATTGGCGTTGCGCAAAAGTTAAACCCAGA

 34750 34760 34770 34780 34790 34800
 AGATCTGGGTGATGATGCTTACAAGAACAGCACGGCGTAAATATGCTTATCATGGCGG

 34810 34820 34830 34840 34850 34860
 TGGCGATGGCAAATGGTATTGCCTCGGTTGAATTGGTTGCGTTAGTAAAGCAGGGCT

 34870 34880 34890 34900 34910 34920
 GTTATGTCATTGGTGTGCAGGTCTAGTGCCTGATGCGGTTGAAGATGCAATTGTCG

 34930 34940 34950 34960 34970 34980
 TATTCAAGCTGAATTACCAATGGCCCTATGCGGTTAACTTGATCCATGCACCAGCAGA

 34990 35000 35010 35020 35030 35040
 AGAAGCATTAGAGCGTGGCGCGGTTGAACGTTTCTAAACTGGCGTAAAGACGGTAGA

 35050 35060 35070 35080 35090 35100
 GGCTTCAGCTTACCTGGTTAACTGAACACATTGTTGGTATCGTGTGCTGGTCTAAC

 35110 35120 35130 35140 35150 35160
 TAAAAACGCAGATGGCAGTGTAAATATCGTAACAAGGTTATCGCTAAAGTATCGCGTAC

 35170 35180 35190 35200 35210 35220
 CGAAGTTGGTCGCCGCTTATGGAACCTGCACCGCAAAATTACTGGATAAGTTATTAGA

 35230 35240 35250 35260 35270 35280
 ACAAAATAAGATCACCCCTGAACAAGCTGCTTAGCGTTGCTTGTACCTATGGCTGATGA

 35290 35300 35310 35320 35330 35340
 TATTACTGGGAAAGCGGATTCTGGTGGTCATAAGATAACCGTCCGTTTAACATTATT

Fig. 5

35350 35360 35370 35380 35390 35400
 ACCGACGATTATTGGTCTGCGTATGAAGTGCAAGCGAAGTATAACTTCTCTGCATT

 35410 35420 35430 35440 35450 35460
 ACGTGTGGTGGTGGTGGTATCGAACGCCTGAAGCAGCACTCGCTGCATTTAACAT

 35470 35480 35490 35500 35510 35520
 GGGCGCGGCTTATATCGTTCTGGGTTCTGTGAATCAGCGTGTGGTGAAGCGGGTGCATC

 35530 35540 35550 35560 35570 35580
 TGAATATACTCGTAAACTGTTATCGACAGTTGAAATGGCTGATGTGACTATGGCACCTGC

 35590 35600 35610 35620 35630 35640
 TGCAGATATGTTGAAATGGGTGTGAAGCTGCAAGTATTAAAACGCCTATGTTCGC

 35650 35660 35670 35680 35690 35700
 GATGCGTGCAGAAGAACTGTATGACTTGTATGACTCGATTGAAGATATCCC

 35710 35720 35730 35740 35750 35760
 AGCTGCTAACGTGAGAAGATTGAAAAACAAATCTTCCGTGCAAACCTAGACGAGATTG

 35770 35780 35790 35800 35810 35820
 GGATGGCACTATCGCTTCTTACTGAACCGCATTGAGAAATGCTAGCCCGTGCAACGAG

 35830 35840 35850 35860 35870 35880
 TAGTCCTAACGTAAAATGGCACTTATCTTCCGTGGTATCTGGCCTTCTTCACGCTG

 35890 35900 35910 35920 35930 35940
 GTCAAACACAGGCAGAGAAGGGACGTGAAATGGATTATCAGATTGGGCAGGCCAAGTTT

 35950 35960 35970 35980 35990 36000
 AGGTGCATTCAACAGCTGGGTGAAAGGTTCTTACCTTGAAGACTATACCCGCCGTGGCGC

 36010 36020 36030 36040 36050 36060
 TGTAGATGTTGCTTGCATATGCTTAAAGGTGCTGCGTATTTACAACGTGAAACCAGTT

 36070 36080 36090 36100 36110 36120
 GAAATTGCAAGGTGTTAGCTTAAGTACAGAATTGGCAAGTTATCGTACGAGTGATTAATG

 36130 36140 36150 36160 36170 36180
 TTACTTGATGATATGTGAATTAAATTAAAGCGCCTGAGGGCGCTTTTGTTTAACT

 36190 36200 36210 36220 36230 36240
 CAGGTGTTGTAACTCGAAATTGCCCTTCAAGTTAGATCGATTACTCACTCACAAATATG

 36250 36260 36270 36280 36290 36300
 TTGATATCGCACTGCCATATACTTGCTCATCCAAAGCCCTATATTGATAATGGTGTAA

 36310 36320 36330 36340 36350 36360
 TAGTCTTAATATCCGAGTCTTCTTCAGCATAATACTAATATAGAGACTCGACCAATGT

 36370 36380 36390 36400 36410 36420
 TAAACACAAACAAAGAATATATTCTGTGACTGCCTTATTATTAACGAGTGCGAGTACGA

 36430 36440 36450 36460 36470 36480
 CAGCTACTACGCTAAACAATTGAGATATCAGCAATTGAACAACGTATTCTGGTCGTATCG

Fig. 5

36490 36500 36510 36520 36530 36540
 GTGTGGCTGTTTAGATACGAAAATAACAAACGTGGGCTTACAATGGTATGCACATT

 36550 36560 36570 36580 36590 36600
 TTCCGATGATGAGTACATTCAAAACCCCTCGCTTGCACGAAATGCTAAGTGAATCGACAA

 36610 36620 36630 36640 36650 36660
 ATGGTAATCTGGATCCCAGTACTAGCTCATTGATAAAGGCTGAAGAATTAATCCCTGGT

 36670 36680 36690 36700 36710 36720
 CACCAAGTCACTAAACGTTGTGAATAACACTATTACAGTGGCGAAAGCGTGTGAAGCAA

 36730 36740 36750 36760 36770 36780
 CAATGCTGACCAAGTGATAATACCGCGGCTAATATTGTTTACAGTATATCGGAGGCCCTC

 36790 36800 36810 36820 36830 36840
 AAGGCAGTTACTGCATTCTGCGAGAAATTGGTATGAAGAGAGTCAGTTAGATCGTATAG

 36850 36860 36870 36880 36890 36900
 AACCTGAATTGAATGAAGCTAAGGTGGAGACTTGCCTGATACACGACACCGAAAGCCA

 36910 36920 36930 36940 36950 36960
 TAGTTACCAACGCTAACAAACTACTACTTGTTGATGTTCTACTTGATTGGATAAAAACC

 36970 36980 36990 37000 37010 37020
 AACTTAAACATGGATGCAAAATAATAAAGTGTCAAGATCCTTACTGCCTCTATATTAC

 37030 37040 37050 37060 37070 37080
 CGCAAGGCTGGTTATTGCCGACCGCTCAGGTGCGGGTGGTAATGGTCTCGAGGTATAA

 37090 37100 37110 37120 37130 37140
 CTGCTATGCTTGGCACTCCGAGCGTCAACCGCTAACATCATCAGTATTATTAACCGAAA

 37150 37160 37170 37180 37190 37200
 CTGAGTTAGCAATGGCAATGCGCAATGAGATTATTGTTGAGATCGGTAAGCTGATATTCA

 37210 37220 37230 37240 37250 37260
 AAGAATACGCGGTGAAATAATAAGTTATTTTGATAATACTTAAACGAGCGTAGCTATC

 37270 37280 37290 37300 37310 37320
 GAAGTGAGGGCGTCAATTAGACACCTTGCTTCCCTACAAATCTAATGTGTATTACCT

 37330 37340 37350 37360 37370 37380
 CGGCTAGTACAATTGCCCTAACGTTATTCTGTCCAGCTTGGCTTAGTGCATTGCGTTA

 37390 37400 37410 37420 37430 37440
 GCCAATGTGAACACCAAGGGACTTGTGTCGTAACCAACTACCAAGCGACTTGTGCGTTT

 37450 37460 37470 37480 37490 37500
 TATCTTTCTTAGACAAACAGAGGTTAAATGAGTGACGCCTTCCAAATCACAGGAATGAA

 37510 37520 37530 37540 37550 37560
 TCCGCATTCAATAAAATCTAACCCGTACCAACTCCGTACAAGTTGATCTTAGTTGTT

 37570 37580 37590 37600 37610 37620
 AAAATCTATAATAATTCAATTACGGAATTAATCCGTACAACGGAGGTTATGGCTAC

Fig. 5

37630 37640 37650 37660 37670 37680
 TGCAAGACTTGATATCCGTTGGATGAAGAAATCAAAGCTAAGGCTGAGAAAGCATCAGC

 37690 37700 37710 37720 37730 37740
 TTTACTCGGCTTAAAAAGTTAACCGAATACGTTGTTGCTTAATGGACGAAGATTCAAC

 37750 37760 37770 37780 37790 37800
 TAAAGTAGTTCTGAGCATGAGAGTATTACCGTTGAAGCGAATGTATTGACCAATTAT

 37810 37820 37830 37840 37850 37860
 GGCTGCTTGTGATGAAGCGAAAGCCCCAAATAAAGCATTACTTGAAAGCCGCTGTATTTAC

 37870 37880 37890 37900 37910 37920
 TCAGAATGGTGAGTTAAGTGAGTTATTCCAAACGTTCAAAGAACTGGATAAAATCAAAA

 37930 37940 37950 37960 37970 37980
 CATGACAGAGCATCTTGACTGTGGCGAAAAAGAGCTAAATGATTTATCCAAACTCAA

 37990 38000 38010 38020 38030 38040
 GCAGCCAAACATATGCAAGCAGGTATTAGCCGCACTCTGGTTTACCTGCTCTGCGCCG

 38050 38060 38070 38080 38090 38100
 TTACCAAAACAAAAATATCCAATTGCTCATTATAGTATCGCGCCAAGCTCAATTAGC

 38110 38120 38130 38140 38150 38160
 CGCGATACTTACCAAGCAATGGCTAAAAAGTTACCACGTTATCCTATCCCTGTTTT

 38170 38180 38190 38200 38210 38220
 CTTTGGCTCAACTTGCCGTCCATAAAGAGTTCATGGGAGTGGTTAGGCAAAGTTAGC

 38230 38240 38250 38260 38270 38280
 TTAATTAAAGCGTTAGAGTACCTTGGGAAATTAACTCTCACATGAGAGCTTACGCCATC

 38290 38300 38310 38320 38330 38340
 GTTGGTTGATTGTTAACTGAACAAGCTGAGTCATTCTACGCTAAATATGGTTCGACGTT

 38350 38360 38370 38380 38390 38400
 CTCTGCGAAATAATGGTCGAGTAAGAATGTCATATCAATGAAAACAGTCAGTTA

 38410 38420 38430 38440 38450 38460
 TTCACTTAACAGTAAGAGTTAGTATAACAGTTGATGAATTAAATTATTATTCGGTA

 38470 38480 38490 38500 38510 38520
 ATCTCATTGCGATCACGCTAGAAGTGCAGCAGCGGGTCAGACCGAGGCCACAATAGCAGCCG

 38530 38540 38550 38560 38570 38580
 TTACGTTAGGGGATGACTTAAAAGATAACTACTACGTCAGTGGCGATCTAGAGGATT

 38590 38600 38610 38620 38630 38640
 AAAGGTTATGATTACAACATTATTATTGTGCTTAATTCTATCCAATATGCGC

 38650 38660 38670 38680 38690 38700
 AAGCTGAAATATCACTGAAGTAGACTTTATGTCAGTGATGATCCCTAAAGATGTTG

 38710 38720 38730 38740 38750 38760
 CCAAATTAAAGATAGGTGAATCCATAACCGAACCTCCAGCCTTATTCTAAGTAACTCATCTA

Fig. 5

38770 38780 38790 38800 38810 38820
 TTCCACTCTCGCGGGAGACGGGTAACATATATTACTCTTCATCAATTGCTAACTTGAAC

 38830 38840 38850 38860 38870 38880
 ATGACTCGATAGAATTGTTATGGCTCAATTGATGGCGAAGATTCCAGCCTTACAAGA

 38890 38900 38910 38920 38930 38940
 TGCTGGTAAATAGCGATAGGTTGTCCTGCTAGTAATGACATCTCCCAGTCCACAGATC

 38950 38960 38970 38980 38990 39000
 TCTATGGCTCGACTTACTCGGCTTATTTCTAATGTTGCGGTATCGATTGAAATTGTG

 39010 39020 39030 39040 39050 39060
 ACTCGCTAACTTTAGAACATGAGCTCGGCCATCTATACGGAGCTGAACATGAAGAAATAT

 39070 39080 39090 39100 39110 39120
 ATGACGACTATGTCTTATGCTGCGATATGTGGAGACTATACGACTATCATGAACTCTA

 39130 39140 39150 39160 39170 39180
 TGCAGCCTGAAATGAAAGAAAACAAATGATAAAGGCATATTCTATTCCCTGAATTAAAAG

 39190 39200 39210 39220 39230 39240
 TGGATGGCTTGCAGTGCAGGAAATGAAAATACGAATAACAAAAGGTTATTTAGACAATA

 39250 39260 39270 39280 39290 39300
 TTGGTCGGTTAGATAGGATTGGGATATTATTCTCATTGGCTCTACTTAGTGCTGTTAT

 39310 39320 39330 39340 39350 39360
 TATGAGTGCCAGTGCTTCTATCTACGATATTGGTCTAACAAAGTATTTATCTATAGACGC

 39370 39380 39390 39400 39410 39420
 TAAGGTGTTATGTATTTAAGGGATGTTCAAGATGAAACTAGGTGAAACGATGTATAGTT

 39430 39440 39450 39460 39470 39480
 GTATAACATTTTCAACGGTTGGAACGTTGATTCTATCGGGTAACAAGACCGCGACGA

 39490 39500 39510 39520 39530 39540
 TCCCGATAAGTCCGATAGTCATTACTTAGTTGGTCAGATGTTAGATGCTGTACTCACG

 39550 39560 39570 39580 39590 39600
 AAGATAATCGGAAATGTGTCAAATAGAAACTGAGCATTGAATATGTGACGTTAGTG

 39610 39620 39630 39640 39650 39660
 AATTAAACCGTGCACGCCAATGCTGAAGGTTACCGTTTGTGTTAGCTTAAGTGG

 39670 39680 39690 39700 39710 39720
 TAGTTGAAAGATTATCCGACTTCAAATGATTATTTCTATAAGTTTCAGAGTTGTA

 39730 39740 39750 39760 39770 39780
 CTATCGATATCTTATAAGTCTAGTGCACAAAACAGAACTATTTATAGCGCTCAAGAAGG

 39790 39800 39810 39820 39830 39840
 CGATAATTGATAATGAATTATGCCCTGTTACTATTAAGAGACTTTAAATGACTGAGAT

 39850 39860 39870 39880 39890 39900
 ATAAGATATGACACGGAAAGAACATATTGATCACAGGCGCAAGTTAGGGTTGGGCCGAGG

Fig. 5

39910 39920 39930 39940 39950 39960
TATGGCCATCGAATTGCAAAATCAGGTATAACTTAGCAGCTTGTGCACGTAGACTTGA
39970 39980 39990 40000 40010 40020
TAATTTAGTTGCACTGAAAGCAGAACACTCTAGCCCTCAATCCTCACATCCAAATCGAAAT
40030 40040 40050 40060 40070 40080
AAAACCTCTTGATGTCAATGAACATGAACAAGTCTTCACTGTTTCCATGAATTCAAAGC
40090 40100 40110 40120 40130
TGAATTGGTACGCTTGATCGTATTATTGTTAATGCTGGATTAGGCAAGGGTGGATCC

Fig. 5

10 20 30 40 50 60
 AAATGCAATTAAATTATGGCGTAAATAGAGTGAAAACATGGCTAATATTCACTAAGTCCTG
 70 80 90 100 110 120
 AATTTATATAAAGTTAATCTGTTATTTAGCGTTACCTGGTCTATCAGTGAGGTT
 130 140 150 160 170 180
 ATAGCCATTATTAGTGGGATTGAAGTGATTAAAGCTATGTATATTATTGCAAATATA
 190 200 210 220 230 240
 AATTGTAACAATTAAGACTTTGGACACTTGAGTTCAATTGAAATTGATTGGCATAAAAT
 250 260 270 280 290 300
 TTAAAACAGCTAAATCTACCTCAATCATTAGCAAATGTATGCAGGTAGATTTTCG
 310 320 330 340 350 360
 CCATTTAAGAGTACACTGTACGCTAGGTTTTGTTAGTGTGCAAATGAACGTTTGAT
 370 380 390 400 410 420
 GAGCATTGTTTTAGAGCACAAAATAGATCCTTACAGGAGCAATAACGCAATGGCTAAA
 430 440 450 460 470 480
 AGAACACCACATCGATTAAGCACGCCAAGGATGTGTTAAGTAGTGTGATCAACAGTTAA
 490 500 510 520 530 540
 ATTCTCGCTTGCAAGAATGTCCGATTGCCATCATTGGTATGGCATCGGTTTGCAGATG
 550 560 570 580 590 600
 CTAAAAACTGGATCAATTCTGGATAACATCGTTGACTCTGTGGACGCTATTATTGATG
 610 620 630 640 650 660
 TGCCTAGCGATCGCTGGAACATTGACGACCATTACTCGGCTGATAAAAAGCAGCTGACA
 670 680 690 700 710 720
 AGACATACTGCAAACCGGGTGGTTCAATTCCAGAGCTTGATTTGATCCGATGGAGTTG
 730 740 750 760 770 780
 GTTTACCGCCAAATATCCTCGAGTTAAGTGCATGGCATTGGTAGTGATTGACCATGATAAAATTGGTA
 790 800 810 820 830 840
 CTCGTGATGTATTAAGTGTGCTGGCATTGGTAGTGATTGACCATGATAAAATTGGTA
 850 860 870 880 890 900
 TCACGCTGGGTGTCGGTGGTCAGAAACAAATTGCCATTAAACGTCGCGCCTACAAG
 910 920 930 940 950 960
 GCCCGGTATTAGAAAAAGTATTAAAAGCCTCAGGCATTGATGAAGATGATCGCGCTATGA
 970 980 990 1000 1010 1020
 TCATCGACAAATTAAAAAGCCTACATCGGCTGGGAAGAGAACTCATTCCCAGGCATGC
 1030 1040 1050 1060 1070 1080
 TAGGTAACGTTATTGCTGGTCGTATGCCAATCGTTTGATTTGGTAGCTAACTGTG
 1090 1100 1110 1120 1130 1140
 TGGTTGATGCGGCATGCGCTGGCTCCCTTGCACTGTTAAAATGGCGATCTCAGACTTAC

Fig. 6

1150 1160 1170 1180 1190 1200
 TTGAATATCGTCAGAAGTCATGATATCGGGTGGTGTATGTTGATAACTCGCCATTCA
 1210 1220 1230 1240 1250 1260
 TGTATATGTCATTCTCGAAAACACCAGCATTTACCACCAATGATGATATCCGTCCGTTG
 1270 1280 1290 1300 1310 1320
 ATGACGATTCAAAAGGCATGCTGGTTGGTGAAGGTATTGGCATGATGGCGTTAACCGTC
 1330 1340 1350 1360 1370 1380
 TTGAAGATGCTGAACGTGACGGCGACAAAATTATTCTGTACTGAAAGGTATCGGTACAT
 1390 1400 1410 1420 1430 1440
 CTTCAGATGGTCGTTCAAATCTATTTACGCTCCACGCCAGATGCCAAGCAAAAGCGC
 1450 1460 1470 1480 1490 1500
 TAAAACGTGCTTATGAAGATGCCGGTTTGCCCCCTGAAACATGTGGTCTAATTGAAGGCC
 1510 1520 1530 1540 1550 1560
 ATGGTACGGGTACCAAAGCGGGTGTGCCCGAGAATTGCTGGCTTGACCAACACTTG
 1570 1580 1590 1600 1610 1620
 GCGCCGCCAGTGTGAAAGCAATATATGCCCTTAGGCTCAGTTAAATCGCAAATTGGTC
 1630 1640 1650 1660 1670 1680
 ATACTAAATCTGCGGCTGGCTCTGCGGGTATGATTAAGCGGGCATTAGCGCTGCATCATA
 1690 1700 1710 1720 1730 1740
 AAATCTTACCTGCAACGATCCATATCGATAAAACCAAGTGAAGCCTGGATATCAAAACA
 1750 1760 1770 1780 1790 1800
 GCCCGTTACCTAAACAGCGAAACCGCTCTGGATGCCACGTGAAGATGGTATTCCAC
 1810 1820 1830 1840 1850 1860
 GTCGTGCAGGTATCAGCTATTGGTTGGCGGGACCAACTTCCATATTATTTAGAAG
 1870 1880 1890 1900 1910 1920
 AGTATGCCCAAGGTACGATAGCGCATATCGCTTAAACTCAGTGAGCCAAACTGTGTTGA
 1930 1940 1950 1960 1970 1980
 TCTCGGCAAACGACCAACAAGGTATTGTTGCTGAGTTAAATACTGGCGTACTAAACTGG
 1990 2000 2010 2020 2030 2040
 CTGTCGATGCTGATCATCAAGGGTTGTATTAATGAGTTAGTGACAACGTGGCCATTAA
 2050 2060 2070 2080 2090 2100
 AAACCCCATCCGTTAACCAAGCTCGTTAGGTTTGTGCGCGTAATGCAAATGAAGCGA
 2110 2120 2130 2140 2150 2160
 TCGCGATGATTGATAACGGCATTGAAACAATTCAATGCGAACCGCAGATAAAATGACATGGT
 2170 2180 2190 2200 2210 2220
 CAGTACCTACCGGGTTACTATCGTCAAGCCGGTATTGATGCAACAGGTAAAGTGGTTG
 2230 2240 2250 2260 2270 2280
 CGCTATTCTCAGGGCAAGGTTCGCAATACGTGAACATGGTCGTGAATTAACTGTAACT

Fig. 6

2290 2300 2310 2320 2330 2340
 TCCCAAGCATGATGCACAGTGCTGCGCGATGGATAAAGAGTTCAAGTGCCTGGTTAG
 2350 2360 2370 * 2380 2390 2400
 GCCAGTTATCTGCAGTTACTTCCCTATCCCTGTTATACGGATGCCGAGCGTAAGCTAC
 2410 2420 2430 2440 2450 2460
 AAGAAGAGCAATTACGTTAACGCAACATGCGCAACCAGCGATGGTAGTTGAGTGTG
 2470 2480 2490 2500 2510 2520
 GTCTGTTCAAAACGTTAACGCAAGCAGGTTAAAGCTGATTTGCTGCCGGTCATAGTT
 2530 2540 2550 2560 2570 2580
 TCGGTGAGTTAACCGCATTATGGGCTGCCGATGTATTGAGCGAAAGCGATTACATGATGT
 2590 2600 2610 2620 2630 2640
 TAGCGCGTAGTCGTGGTCAAGCAATGGCTGCCAGAGCAACAAGATTTGATGCAGGTA
 2650 2660 2670 2680 2690 2700
 AGATGGCCGCTGTTGGTGAATCAAAGCAAGTCGCTGTGATCATTGATACCCCTGATG
 2710 2720 2730 2740 2750 2760
 ATGTCTCTATTGCTAACTCAACTCGAATAACCAAGTTGTTATTGCTGGTACTACGGAGC
 2770 2780 2790 2800 2810 2820
 AGGTTGCTGTAGCGGTTACAACCTTAGGTAATGCTGGTTCAAAGTTGTGCCACTGCCGG
 2830 2840 2850 2860 2870 2880
 TATCTGCTGCGTCCATACACCTTAGTTGTCACCGCAAAACCATTTGCTAAAGCGG
 2890 2900 2910 2920 2930 2940
 TTGATAGCGCTAAATTAAAGCGCCAAGCATTCCAGTGTGTTGCTAATGGCACAGGCTGG
 2950 2960 2970 2980 2990 3000
 TGCATTCAAGCAAACCGAATGACATTAAGAAAAACCTGAAAAACACATGCTGGAATCTG
 3010 3020 3030 3040 3050 3060
 TTTCATTTCAATCAAGAAATTGACAACATCTATGCTGATGGTGGCCCGTATTTATCGAAT
 3070 3080 3090 3100 3110 3120
 TTGGTCAAAGAATGTATTAACCAAATTGGTTGAAACATTCTCACTGAAAAATCTGATG
 3130 3140 3150 3160 3170 3180
 TGACTGCTATCGCGTTAATGCTAACCTAAACACCGTACAAATGCGCCAAG
 3190 3200 3210 3220 3230 3240
 CTGCGCTGCAAATGGCAGTGCTTGGTGTGCGATTAGACAATATTGACCCGTACGACGCCG
 3250 3260 3270 3280 3290 3300
 TTAAGCGTCCACTTGTGCGCCGAAAGCATCACCAATGTTGATGAAGTTATCTGCAGCGT
 3310 3320 3330 3340 3350 3360
 CTTATGTTAGTCCGAAACGAAGAAAGCGTTGCTGATGCATTGACTGATGGCTGGACTG
 3370 3380 3390 3400 3410 3420
 TTAAGCAAGCGAAAGCTGTACCTGCTGTTGTCACAACCACAAGTGATTGAAAAGATCG

Fig. 6

3430 3440 3450 3460 3470 3480
 TTGAAGTTGAAAAGATAGTTGAACGCATTGTCGAAGTAGAGCGTATTGTCGAAGTAGAAA
 3490 3500 3510 * 3520 3530 3540
 AAATCGTCTACGTTAATGCTGACGGTTCGCTTATATCGAAAATAATCAAGACGTTAAC
 3550 3560 3570 3580 3590 3600
 GCGCTGTTAGCAACGTGACTAATAGCTCAGTGACTCATAGCAGTGATGCTGACCTTG
 3610 3620 3630 3640 3650 3660
 TTGCCTCTATTGAACCGAGTGTGGTCAATTGTTGCACACCAACAGCAATTATTAAATG
 3670 3680 3690 3700 3710 3720
 TACATGAACAGTTATGCAAGGTCCACAAGACTACGCGAAAACAGTGCAGAACAGTACTTG
 3730 3740 3750 3760 3770 3780
 CTGCGCAGACGAGCAATGAATTACCGGAAAGTTAGACCGTACATTGTCTATGTATAACG
 3790 3800 3810 3820 3830 3840
 AGTTCCAATCAGAAACGCTACGTGTACATGAAACGTACCTGAACAATCAGACGAGCAACA
 3850 3860 3870 3880 3890 3900
 TGAACACCATGTTACTGGTGCTGAAGCTGATGTGCTAGCAACCCATAACTCAGGTAG
 3910 3920 3930 3940 3950 3960
 TGAATACAGCCGTTGCCACTAGTCACAAGGTAGTTGCTCCAGTTATTGCTAATACAGTGA
 3970 3980 3990 4000 4010 4020
 CGAATGTTGATCTAGTGTCAAGTAACGGCGGGTTGCAGTGCAAACACTGTGGCATTAG
 4030 4040 4050 4060 4070 4080
 CGCCTACGCAAGAAATCGCTCCAACAGTCGCTACTACGCCAGCACCCGCATTGGTTGCTA
 4090 4100 4110 4120 4130 4140
 TCGTGGCTGAACCTGTGATTGTTGCGCATGTTGCTACAGAAGTTGCACCAATTACACCAT
 4150 4160 4170 4180 4190 4200
 CAGTTACACCAGTTGTCGCAACTCAAGCGGCTATCGATGTAGCAACTATTAACAAAGTAA
 4210 4220 4230 4240 4250 4260
 TGTAGAAGTTGCTGATAAAACCGGTTATCCAACGGATATGCTGGAACGTGAGCATGG
 4270 4280 4290 4300 4310 4320
 ACATGGAAGCTGACTTAGGTATCGACTCAATCAAACGTGTTGAGATATTAGGCGCAGTAC
 4330 4340 4350 4360 4370 4380
 AGGAATTGATCCCTGACTTACCTGAACCTAACCTGAAAGATCTTGCTGAGCTACGCACGC
 4390 4400 4410 4420 4430 4440
 TTGGTGAGATTGTCGATTACATGAATTCAAAGCCCAGGCTGTAGCTCCTACAAACAGTAC
 4450 4460 4470 4480 4490 4500
 CTGTAACAAGTGCACCTGTTGCCTGCATCTGCTGGTATTGATTAGCCCACATCCAAA
 4510 4520 4530 4540 4550 4560
 ACGTAATGTTAGAAGTGGTTGCAGACAAACCGGTTACCCAACAGACATGCTAGAACTGA

Fig. 6

4570 4580 4590 4600 4610 4620
 GCATGGATATGGAAGCTGACTTAGGTATTGATTCAATCAAGCGTGTGGAAATCTTAGGTG
 4630 4640 4650 4660 4670 4680
 CAGTACAGGAGATCATAACTGATTACCTGAGCTAAACCCCTGAAGATCTTGCTGAATTAC
 4690 4700 4710 4720 4730 4740
 GCACCCTAGGTGAAATCGTTAGTTACATGCAAAGCAAAGCGCCAGTCGCTGAAAGTGC
 4750 4760 4770 4780 4790 4800
 CAGTGGCGACGGCTCCTGTAGCAACAAAGCTCAGCACCGTCTATCGATTGAACCACATTC
 4810 4820 4830 4840 4850 4860
 AAACAGTGATGATGGATGTAGTTGCAGATAAGACTGGTTATCCAACGTGACATGCTAGAAC
 4870 4880 4890 4900 4910 4920
 TTGGCATGGACATGGAAGCTGATTAGGTATCGATTCAATCAAACGTGTGGAAATATTAG
 4930 4940 4950 4960 4970 4980
 GCGCAGTGCAAGGAGATCATCACTGATTACCTGAGCTAAACCCAGAAGACCTCGCTGAAT
 4990 5000 5010 5020 5030 5040
 TACGCACGCTAGGTGAAATCGTTAGTTACATGCAAAGCAAAGCGCCAGTCGCTGAGAGTG
 5050 5060 5070 5080 5090 5100
 CGCCAGTAGCGACGGCTCTGTAGCAACAAAGCTCTGCACCGTCTATCGATTAAACCATA
 5110 5120 5130 5140 5150 5160
 TCCAAACAGTGATGATGGAAGTGGTTGCAGACAAAACCGGTTATCCAGTAGACATGTTAG
 5170 5180 5190 5200 5210 5220
 AACTTGCTATGGACATGGAAGCTGACCTAGGTATCGATTCAATCAAGCGTGTAGAAATT
 5230 5240 5250 5260 5270 5280
 TAGGTGCGGTACAGGAAATCATTACTGACTTACCTGAGCTAACCTGAAGATCTTGCTG
 5290 5300 5310 5320 5330 5340
 AACTACGTACATTAGGTGAAATCGTTAGTTACATGCAAAGCAAAGCGCCCGTAGCTGAAG
 5350 5360 5370 5380 5390 5400
 CGCCTGCAGTACCTGTTGCAGTAGAAAGTGCACCTACTAGTGTAAACAAGCTCAGCACCGT
 5410 5420 5430 5440 5450 5460
 CTATCGATTTAGACCACATCCAAAATGTAATGATGGATGTTGCTGATAAGACTGGTT
 5470 5480 5490 5500 5510 5520
 ATCCTGCCAATATGCTTGAATTAGCAATGGACATGGAAGCCGACCTTGGTATTGATTCAA
 5530 5540 5550 5560 5570 5580
 TCAAGCGTGTGAAATTCTAGGCGCGGTACAGGAGATCATTACTGATTACCTGAACCAA
 5590 5600 5610 5620 5630 5640
 ACCCAGAAGACTTAGCTGAACCTACGTACGTTAGAAGAAATTGTAACCTACATGCAAAGCA
 5650 5660 5670 5680 5690 5700
 AGGCGAGTGGTGTACTGTAATGTAGTGGCTAGCCCTGAAAATAATGCTGTATCAGATG

Fig. 6

5710 5720 5730 5740 5750 5760
 CATTATGAAAGCAATGTGGCGACTATCACAGCGGCCGAGAACATAAGGCGGAATTAA
 5770 5780 5790 5800 5810 5820
 AACCGGCCGAGCGAACCGTTGCTATCTCTCGTCTAAGCTCTATCAGTAAAATAAGCC
 5830 5840 5850 5860 5870 5880
 AAGATTGTAAGGTGCTAACGCCCTAACGCTAGCTAGTGGACTGATAATGCTGTGTTAC
 5890 5900 5910 5920 5930 5940
 TTGCAGACCACCTATTGCAAACCTGGCTGGAATGTAACGCATTGCAACCAACTGGGTAG
 5950 5960 5970 5980 5990 6000
 CTGTAACAAACGACGAAAGCATTAAATAAGTCAGTGAACCTGGTGAACCTTAAATGGCGTTG
 6010 6020 6030 6040 6050 6060
 ATGAAACTGAAATCAACACATTATTACTGCTAACGCACAATTGGATGCAGTTATCTATC
 6070 6080 6090 6100 6110 6120
 TGCACGCAAGTAGCGAAATTAAATGCTATCGAATACCCACAAGCATAAGCAAGGCCTGA
 6130 6140 6150 6160 6170 6180
 TGTTAGCCTTCTTATTAGCGAAATTGAGTAAAGTAACCTAACGCCGCTAAAGTGCCTGGCG
 6190 6200 6210 6220 6230 6240
 CCTTTATGATTGTTACTCAGCAGGGTGGTCATTAGGTTTGATGATATCGATTCTGCTA
 6250 6260 6270 6280 6290 6300
 CAAGTCATGATGTGAAAACAGACCTAGTACAAAGCGGCTAACCGTTAGTTAAGACAC
 6310 6320 6330 6340 6350 6360
 TGTCTCACGAGTGGGATAACGTATTCTGTCGTGCGGGTGTGATATTGCTTCGTCAATTACGG
 6370 6380 6390 6400 6410 6420
 CTGAACAAGTTGCAAGCCTTGTAGTGTGAACTACTTGATGCTAACACTGTATTAACAG
 6430 6440 6450 6460 6470 6480
 AAGTGGTTATCAACAAGCTGGTAAAGGCCCTGAACGTATCACGTTAACCTGGTGTGGCTA
 6490 6500 6510 6520 6530 6540
 CTGACAGCTATGCATTAACAGCTGGCAATAACATCGATGCTAACCGGTATTTAGTGA
 6550 6560 6570 6580 6590 6600
 GTGGTGGCGAAAAGGTGTAAGTGCACATTGTGTTGCTCGTATAGCTAAAGAATATCAGT
 6610 6620 6630 6640 6650 6660
 CTAAGTTCATCTTATTGGGACGTTAACGTTCTCAAGTGACGAACCGAGCTGGCAAGTG
 6670 6680 6690 6700 6710 6720
 GTATTACTGATGAAGCGCGTTAAAGAAAGCAGCGATGCAGTCTTGTGATTACAGCAGGTG
 6730 6740 6750 6760 6770 6780
 ATAAACCAACACCCGTTAACGATCGTACAGCTAACCAACCAAGCTAACCGTAAACGTGAAA
 6790 6800 6810 6820 6830 6840
 TTGCGAACCTTGTCTGCAATTACCGCTGCTGGTGGCCAAGCTGAATATGTTCTGCAG

Fig. 6

6850 6860 6870 6880 6890 6900
 ATGTAACATAATGCAGCAAGCGTACAAATGGCAGTCGCTCCAGCTATCGCTAAGTTCGGTG
 6910 6920 6930 6940 6950 6960
 CAATCACTGGCATCATTGAGCTGGCGGGGTGTTAGCTGACCAATTGAGCAAAAAAA
 6970 6980 6990 7000 7010 7020
 CACTGAGTGATTTGAGTCTGTTACAGCACTAAAATTGACGGTTGTTATCGCTACTAT
 7030 7040 7050 7060 7070 7080
 CAGTCACTGAAGCAAGCAACATCAAGCAATTGGTATTGTTCTCGTCAGCGGCTGGTTCT
 7090 7100 7110 7120 7130 7140
 ACGGTAACCCCGGCCAGTCTGATTACTCGATTGCCAATGAGATCTAAATAAAACCGCAT
 7150 7160 7170 7180 7190 7200
 ACCGCTTAAATCATTGCACCCACAAGCTCAAGTATTGAGCTTAACTGGGTCTTGGG
 7210 7220 7230 7240 7250 7260
 ACGGTGGCATGGTAACGCCCTGAGCTAACGTATGTTGACCAACGTGGTGTACATTA
 7270 7280 7290 7300 7310 7320
 TTCCACTTGATGCAGGTGCACAGTTATTGCTGAATGAACTAGCCGCTAATGATAACCGTT
 7330 7340 7350 7360 7370 7380
 GTCCACAAATCCTCGTGGTAATGACTTATCTAAAGATGCTAGCTCTGATCAAAAGTCTG
 7390 7400 7410 7420 7430 7440
 ATGAAAAGAGTACTGCTGTAAAAAGCCACAAGTTAGTCGTTATCAGATGCTTAGTAA
 7450 7460 7470 7480 7490 7500
 CTAAAAGTATCAAAGCGACTAACAGTAGCTCTTATCAAACAAAGACTAGTGCTTATCAG
 7510 7520 7530 7540 7550 7560
 ACAGTAGTGCTTTCAAGGTTAACGAAAACCACTTTAGCTGACCACATGATCAAAGGCA
 7570 7580 7590 7600 7610 7620
 ATCAGGTATTACCAACGGTATGCGCGATTGCTGGATGAGTGATGCAGCAAAGCGACTT
 7630 7640 7650 7660 7670 7680
 ATAGTAACCGAGACTGTGCATTGAAGTATGTCGGTTCGAAGACTATAATTGTTAAAG
 7690 7700 7710 7720 7730 7740
 GTGTGGTTTTGATGGCAATGAGGCAGCGGATTACCAATTGTCGCGCTGTGACAA
 7750 7760 7770 7780 7790 7800
 GGGCGTCAGAACAGGATTCTGAAGTCCGTATTGCCCAAAGATCTTAGCCTGAAAGTG
 7810 7820 7830 7840 7850 7860
 ACGGTAAACCTGTGTTCATTATGCAGCGACAATATTGTTAGCAACTCAGCCACTTAATG
 7870 7880 7890 7900 7910 7920
 CTGTGAAGGTAGAACTTCCGACATTGACAGAAAGTGTGATAGCAACAAATAAGTAACG
 7930 7940 7950 7960 7970 7980
 ATGAAGCACAAAGCGTTACAGCAATGGCACCTTGTCCACGGTAAAGTCTGCAGGGCA

Fig. 6

7990 8000 8010 8020 8030 8040
 TTAAGCAGATATTAAGTTGTGACGACAAGGGCTGCTATTGGCTTGTCAAGATAACCGATG

 8050 8060 8070 8080 8090 8100
 TTGCAACAGCTAACGCAAGGATCCTCCCGTTAGCTGACAACAATATCTTGCCAATGATT

 8110 8120 8130 8140 8150 8160
 TGGTTTATCAGGCTATGTTGGTCTGGGTGCGCAAACAAATTGGTTAGGTAGCTTACCTT

 8170 8180 8190 8200 8210 8220
 CGGTGACAACGGCTTGGACTGTGTATCGTGAAGTGGTTGTAGATGAAGTATTTATCTGC

 8230 8240 8250 8260 8270 8280
 AACTTAATGTTGTTGAGCATGATCTATTGGGTCACGCCAGTAAAGCCGTTGTGATA

 8290 8300 8310 8320 8330 8340
 TTCAATTGATTGCTGCTGATATGCAATTACTGCCGAAGTGAAATCAGCGCAAGTCAGTG

 8350 8360 8370 8380 8390 8400
 TCAGTGACATTTAACGATATGTCATGATCGAGTAAATAAACGATAGGCATGGTGT

 8410 8420 8430 8440 8450 8460
 GAGCATGGCGTCTGCTTCTTCATTTAACATTAACAATTAGCTAAACGGT

 8470 8480 8490 8500 8510 8520
 TGCTTAAACCAAGTAAACAAGTGCTTTAGCTATTACTATTCAAACAGGATATTAAAG

 8530 8540 8550 8560 8570 8580
 AGAATATGACGGAATTAGCTGTTATTGGTATGGATGCTAAATTAGCGGACAAGACAATA

 8590 8600 8610 8620 8630 8640
 TTGACCGTGTGGAACCGCGCTTCTATGAAGGTGCTTATGTAGGTAATGTTAGCCCGTAA

 8650 8660 8670 8680 8690 8700
 GTACCGAATCTAATGTTATTAGCAATGGCGAAGAACAGTTATTACTGCCATGACAGTT

 8710 8720 8730 8740 8750 8760
 TTAACCTCTGTCAGTCTACTAGCGCAAACGAATCAGTTAAATATAGCTGATATCGCGGT

 8770 8780 8790 8800 8810 8820
 TGCTGATTGCTGATGTAAAAAGTGCTGATGATCAGCTTGTAGTCCAAATTGCATCAGCAA

 8830 8840 8850 8860 8870 8880
 TTGAAAAACAGTGTGCGAGTTGTGTTATTGCTGATTTAGGCCAAGCATTAAATCAAG

 8890 8900 8910 8920 8930 8940
 TAGCTGATTTAGTTAATAACCAAGACTGTCCTGTGGCTGTAATTGGCATGAATAACTCGG

 8950 8960 8970 8980 8990 9000
 TTAATTATCTCGTCATGATCTTGAATCTGTAAGTGCAACAACTCAGCTTGATGAAACCT

 9010 9020 9030 9040 9050 9060
 TCAATGGTTATAACAATGTAGCTGGGTCGCGAGTTACTTATCGCTTCAACTGCGTT

 9070 9080 9090 9100 9110 9120
 CCAATGCTAACGCAATGTTATATACGCCAACATTAAGGGCTCGCTCAATCGGGCGTAA

Fig. 6

9130 9140 9150 9160 9170 9180
 ATGCTCAATTAAACGTTGGAAACATTAGCGATACTGCAAAGACCGCATTGCAGCAAGCTA

 9190 9200 9210 9220 9230 9240
 GCATAACTGCAGAGCAGGTTGGTTGTTAGAAGTGTCAAGCAGTCGCTGATTGGCAATCG

 9250 9260 9270 9280 9290 9300
 CATTGTCTGAAAGCCAAGGTTAACATGTCTGCTTATCATACGCAAACCTTGCATACTG

 9310 9320 9330 9340 9350 9360
 CATTAAGCAGTGCCCCGTAGTGTGACTGGTGAAGGCGGGTGTTTCACAGGTCGCAGGTT

 9370 9380 9390 9400 9410 9420
 TATTGAAATGTGTAATTGGTTACATCAACGTTATATTCCGGCGATTAAAGATTGGCAAC

 9430 9440 9450 9460 9470 9480
 AACCGAGTGACAATCAAATGTCACGGTGGCGGAATTCAACCATTCTATATGCCTGTAGATG

 9490 9500 9510 9520 9530 9540
 CTCGACCTTGGTTCCCACATGCTGATGGCTCTGCACACATTGCCGCTTATAGTTGTGTGA

 9550 9560 9570 9580 9590 9600
 CTGCTGACAGCTATTGTCATATTCTTTACAAGAAAACGTCTTACAAGAACTGTGTTGA

 9610 9620 9630 9640 9650 9660
 AAGAAACAGTCTTGCAGATAATGACTTAACGAAAGCAAGCTTCAGACTCTTGAACAAA

 9670 9680 9690 9700 9710 9720
 ACAATCCAGTAGCTGATCTGCGCACTAATGGTTACTTGCATCGAGCGAGTTAGCATTAA

 9730 9740 9750 9760 9770 9780
 TCATAGTACAAGGTAATGACGAAGCACAATTACGCTGTGAATTAGAAAATTACAGGGC

 9790 9800 9810 9820 9830 9840
 AGTTAAGTACTACTGGCATAAGTACTATCAGTATTAAACAGATCGCAGCAGACTGTTATG

 9850 9860 9870 9880 9890 9900
 CCCGTAATGATACTAACAAAGCCTATAGCGCAGTGCTTATTGCCGAGACTGCTGAAGAGT

 9910 9920 9930 9940 9950 9960
 TAAGCAAAGAAAATAACCTTGGCGTTGCTGGTATCGCTAGCGTGTAAATGAAGATGCTA

 9970 9980 9990 10000 10010 10020
 AAGAATGGAAAACCCCGAAGGGCAGTTATTTACCGCGCAGCCTGCAAATAACAGGCTG

 10030 10040 10050 10060 10070 10080
 CTAACAGCACACAGAATGGTGTACCTTCATGTACCCAGGTATTGGTGTACATATGTTG

 10090 10100 10110 10120 10130 10140
 GTTTAGGGCGTGATCTATTCTATTCCCACAGATTATCAGCCTGTAGCGGCTTTAG

 10150 10160 10170 10180 10190 10200
 CCGATGACATTGGCGAAAGTCTAAAGATACTTACTTAATCCACGCGAGTATTAGTCGTC

 10210 10220 10230 10240 10250 10260
 ATAGCTTAAAGAACTCAAGCAGTTGGATCTGGACCTGCGCGGTAACTTAGCCAATATCG

Fig. b

10270 10280 10290 10300 10310 10320
 CTGAAGCCGGTGTGGGTTTGCTTGTGTACCAAGGTATTTGAAGAAGTCTTGCCG

 10330 10340 10350 10360 10370 10380
 TTAAAGCTGACTTGTACAGGTATAGCATGGGTGAAGTAAGCATGTATGCAGCACTAG

 10390 10400 10410 10420 10430 10440
 GCTGCTGGCAGCAACCGGATTGATGAGTGCTCGCCTGCACAATCGAACATCTTAATC

 10450 10460 10470 10480 10490 10500
 ATCAAACTTGCGGCAGTTAAGAACACTACGTCAGCATTGGGCATGGATGATGTAGCTA

 10510 10520 10530 10540 10550 10560
 ACGGTACGTTCGAGCAGATCTGGAAACCTATACCATTAAAGGCAACGATTGAACAGGTCG

 10570 10580 10590 10600 10610 10620
 AAATTGCCTCTGCAGATGAAGATCGTGTATTGCACCATTATCAATACACCTGATAGCT

 10630 10640 10650 10660 10670 10680
 TGTTGTTAGCCGGTTATCCAGAAGCCTGTCAGCAGTCATTAAAGAATTAGGTGTGCGTG

 10690 10700 10710 10720 10730 10740
 CAATGGCATTGAATATGGCGAACGCAATTCACAGCGGCCAGCTTATGCCGAATACGATC

 10750 10760 10770 10780 10790 10800
 ATATGGTTGAGCTATACCATATGGATGTTACTCCACGTATTAAACCAAGATGTATTCAA

 10810 10820 10830 10840 10850 10860
 GCTCATGTTATTTACCGATTCCACAAACGCAGCAAAGCGATTCCCACAGTATTGCTAAAT

 10870 10880 10890 10900 10910 10920
 GTTTGTGTGATGTTGGATTCCACGTTGGTTAACCTTACATGACAAAGGTGCGC

 10930 10940 10950 10960 10970 10980
 GGGTATTCAATTGAAATGGGTCCAGGTCGTTATGTAGCTGGTAGATAAGATCTTAG

 10990 11000 11010 11020 11030 11040
 TTAATGGCGATGGCGATAATAAAAGCAAAGCCAACATGTATCTGTTCTGTGAATGCCA

 11050 11060 11070 11080 11090 11100
 AAGGCACCAGTGTAACTTACTTATTCGTGCGATTGCTAAGTTAATTAGTCATGGCG

 11110 11120 11130 11140 11150 11160
 TGAATTGAAATTAGATAGCTTGTAAACGGGTCATCCTGGTTAAAGCAGGCCATATAG

 11170 11180 11190 11200 11210 11220
 CAAACACGAACAAATAGTCACATCGATATCTAGCGCTGGTGAGTTACCTCATTAGTT

 11230 11240 11250 11260 11270 11280
 GAAATATGGATTAAAGAGAGTAATTATGGAAAATATTGCAGTAGTAGGTATTGCTAATT

 11290 11300 11310 11320 11330 11340
 TGTTCCGGGCTCACAAAGCACCGATCAATTGGCAGCAATTGCTTGAACAACAAGATT

 11350 11360 11370 11380 11390 11400
 GCCGCAGTAAGGCACCGCTGTTCAAATGGCGTTGATCCTGCTAAATATACCGCCAACA

Fig. 6

11410 11420 11430 11440 11450 11460
 AAGGTGACACAGATAAAATTTACTGTGTGCACGGCGGTTACATCAGTGATTCAATTG

 11470 11480 11490 11500 11510 11520
 ATGCTTCAGGTTATCAACTCGATAATGATTATTTAGCCGGTTAGATGACCTTAATCAAT

 11530 11540 11550 11560 11570 11580
 GGGGGCTTATGTTACGAAACAAGCCCTTACCGATGCGGGTTATTGGGGCAGTACTGCAC

 11590 11600 11610 11620 11630 11640
 TAGAAAATCTGTTGATTTAGGTAAATTGTCATTCCAACTAAATCATCTAATCAGC

 11650 11660 11670 11680 11690 11700
 TGTTTATGCCTTGTATCATCAAGTTGTTGATAATGCCCTAAAGGCGGTATTACATCCTG

 11710 11720 11730 11740 11750 11760
 ATTTTCAATTAAACGCATTACACAGCACCGAAAAAAACACATGCTGACAATGCATTAGTAG

 11770 11780 11790 11800 11810 11820
 CAGGTTATCCAGCTGCATTGATCGCGCAAGCGGGCTTGGTGGTCACATTTGCAC

 11830 11840 11850 11860 11870 11880
 TGGATGCGGCTTGTGCTTCATCTGTTAGCGTTAAGTTAGCGTGTGATTACCTGCATA

 11890 11900 11910 11920 11930 11940
 CGGGTAAAGCCAACATGATGCTTGCTGGTGGTATCTGCAGCAGATCCTATGTTCGTAA

 11950 11960 11970 11980 11990 12000
 ATATGGGTTCTCGATATTCAAGCTAACCCAGCTAACAAATGTACATGCCCGTTGACC

 12010 12020 12030 12040 12050 12060
 AAAATTCAAAAGGCTTATTCGCCGGTGAAGGCGCGGCATGATGGTATTGAAACGTCAAA

 12070 12080 12090 12100 12110 12120
 GTGATGCAGTACGTGATGGTGATCATATTACGCCATTATTAAAGGCGGCATTATCGA

 12130 12140 12150 12160 12170 12180
 ATGACGGTAAAGGCGAGTTGTATTAAGCCCGAACACCAAGGGCCAAGTATTAGTATATG

 12190 12200 12210 12220 12230 12240
 AACGTGCTTATGCCGATGCAGATGTTGACCCGAGTACAGTTGACTATATTGAATGTCATG

 12250 12260 12270 12280 12290 12300
 CAACGGGCACACCTAACGGTGACAATGTTGAATTGCGTCGATGGAAACCTTTTCAGTC

 12310 12320 12330 12340 12350 12360
 GCGTAAATAACAAACCATTACTGGGCTCGGTTAAATCTAACCTGGTCATTGTTAACTG

 12370 12380 12390 12400 12410 12420
 CCGCTGGTATGCCTGGCATGACCAAGCTATGTTAGCGCTAGGTAAAGGTCTTATTCCCTG

 12430 12440 12450 12460 12470 12480
 CAACGATTAACCTAACGCAACCCTGCAATCTAAACACGGTTACTTTACTGGCGAGCAAA

 12490 12500 12510 12520 12530 12540
 TGCCAACGACGACTGTGTCTTGGCCAACAACTCCGGGTGCCAAGGCAGATAAACCGCGTA

Fig. 6

12550 12560 12570 12580 12590 12600
 CCGCAGGTGTGAGCGTATTTGGTTGGCAGCAACGCCATTGGTATTACAACAGC

 12610 12620 12630 12640 12650 12660
 CAACGCAAACACTCGAGACTAATTTAGTGTGCTAACACAGTGAGCCTTGGCTATTA

 12670 12680 12690 12700 12710 12720
 TTGGTATGGACAGCCATTTGGTAGTGCCAGTAATTAGCGCAGTTCAAAACCTTATTAA

 12730 12740 12750 12760 12770 12780
 ATAATAATCAAACACCTTCCGTGAATTACCAAGAACACGCTGGAAAGGCATGGAAAGTA

 12790 12800 12810 12820 12830 12840
 ACGCTAACGTCATGCAGTCGTTACAATTACGCAAAGCGCCTAAAGGCAGTTACGTTGAAC

 12850 12860 12870 12880 12890 12900
 AGCTAGATATTGATTTCTTGCCTTAAAGTACCGCCTAATGAAAAAGATTGCTTGATCC

 12910 12920 12930 12940 12950 12960
 CGCAACAGTTAATGATGATGCAAGTGGCAGACAATGCTGCGAAAGACGGAGGTCTAGTTG

 12970 12980 12990 13000 13010 13020
 AAGGTCGTAATGTTGCGGTATTAGTAGCGATGGCATGGAACTGGAATTACATCAGTATC

 13030 13040 13050 13060 13070 13080
 GTGGTCCGCTTAATCTAACCAACCCAAATTGAAGACAGCTTATTACAGCAAGGTATTAACC

 13090 13100 13110 13120 13130 13140
 TGACTGTTGAGCAACGTGAAGAACTGACCAATATTGCTAAAGACGGTGTGCCTCGGCTG

 13150 13160 13170 13180 13190 13200
 CACAGCTAAATCAGTATACGAGTTCTGGTAATATTATGGCGTCACGTATTCGGCGT

 13210 13220 13230 13240 13250 13260
 TATGGGATTTCTGGCCTGCTATTACCGTATCGGCTGAAGAAAATCTGTTATCGTT

 13270 13280 13290 13300 13310 13320
 GTGTTGAATTAGCTGAAAATCTATTCAAACCCAGTGATGTTGAAGCCGTTATTATTGCTG

 13330 13340 13350 13360 13370 13380
 CTGTTGATTTGTCGGTCAATTGAAAACATTACTTACGTCAAGCACTACGGTCCAGTTA

 13390 13400 13410 13420 13430 13440
 ATGAAAAGGGATCTGTAAGTGAATGTGGTCCGGTTAATGAAAGCAGTCAGTAACCAACA

 13450 13460 13470 13480 13490 13500
 ATATTCTGATCAGCAACAATGGCTGGTGGGTGAAGGCGCAGCGGCTATTGTCGTTAAC

 13510 13520 13530 13540 13550 13560
 CGTCATCGCAAGTCACTGCTGAGCAAGTTATGCCGTATTGATGCGGTGAGTTGCCC

 13570 13580 13590 13600 13610 13620
 CTGGTAGCAATGCGAAAGCAATTACGATTGCAGCGGATAAGCATTAAACACTTGCTGGTA

 13630 13640 13650 13660 13670 13680
 TCAGTGCTGCTGATGTAGCTAGTGTGAAAGCACATGCAAGTGGTTAGTGCCGAAATA

Fig. 6

13690 13700 13710 13720 13730 13740
 ATGCTGAAAAACCGCGTTACCGACTTATACCCAAAGCGCAAGTATCAGTCGGTGAAAG

 13750 13760 13770 13780 13790 13800
 CCAATATTGGTCATACGTTAACGCTCGGGTATGGCGAGTATTATTAAAACGGCGCTGC

 13810 13820 13830 13840 13850 13860
 TGTTAGATCAGAATACGAGTCAGAAAAGCAAACATATTGCTATTACGGTCTAG

 13870 13880 13890 13900 13910 13920
 GTCGTGATAACAGCTGCGCGATCTTATCTTATCGAGTTCAAGCGCAAGCGCATCAAGTTG

 13930 13940 13950 13960 13970 13980
 CACCAGCGCCCTGTATCTGGTATGGCAAGCAACGCCACAGTTAGTTAAAACCATCAAAC

 13990 14000 14010 14020 14030 14040
 TCGGTGGTCAGTTAATTAGCAACCGGATTGTTAACAGTGCAGGTTCATCTTACACGCTA

 14050 14060 14070 14080 14090 14100
 TTAAAGCGCAGTTGCCGGTAAGCACTTAAACAAAGTTAACAGCCAGTGTGATGGATA

 14110 14120 14130 14140 14150 14160
 ACCTGAAGCCCCAAGGTATTAGCGCTCATGCAACCAATGAGTATGTGGTGAUTGGAGCTG

 14170 14180 14190 14200 14210 14220
 CTAACACTCAAGCTTCTAACATTCAAGCATCTCATGTTCAAGCGTCAAGTCATGCACAAG

 14230 14240 14250 14260 14270 14280
 AGATAGCACCAACCAAGTTCAAAATATGCAAGCTACAGCAGCCGCTGTAAGTTCACCCC

 14290 14300 14310 14320 14330 14340
 TTCTCAACATCAACACACAGCGCAGCCCGTAGCGGCACCGAGCGTTGGAGTGACTG

 14350 14360 14370 14380 14390 14400
 TGAAACATAAGCAAGTAACCAAATTCACTCAGCAAGCGTCTACGCATAAAGCATTTCAG

 14410 14420 14430 14440 14450 14460
 AAAGTCGTTAGCTGCACAGAAAAACCTATCGCAACTTGTGAATTGCAAACCAAGCTGT

 14470 14480 14490 14500 14510 14520
 CAATCCAAACTGGTAGTGACAATACATCTAACAAACTGCGTCAACAAGCAATACTG

 14530 14540 14550 14560 14570 14580
 TAAACAAATCCTGTATCAGAACGCCATTAACACTTGTGTCTATGCGCCTGTAGTACCGA

 14590 14600 14610 14620 14630 14640
 CAAACCTAACCAAGTACAGAAGCAAAAGCGCAAGCAGCTGCTACACAAGCTGGTTTCAGA

 14650 14660 14670 14680 14690 14700
 TAAAAGGACCTGTTGGTTACAACATATCCACCGCTGCAGTTAATTGAACGTTATAATAAAC

 14710 14720 14730 14740 14750 14760
 CAGAAAACGTGATTACGATCAAGCTGATTGGTTGAATTCGCTGAAGGTGATATTGGTA

 14770 14780 14790 14800 14810 14820
 AGGTATTGGTGCTGAATAACATATTATTGATGGCTATTGCGCGTGTACGTCTGCCAA

Fig. 6

14830 14840 14850 14860 14870 14880
 CCTCAGATTACTTGTAGAACACACGTGTTACTGAACCTGATGCCAAGGTGCATGAATACA

 14890 14900 14910 14920 14930 14940
 AGAAATCATACATGTGTAATGATGTGCCTGTTGATGCACCGTTCTTAATTGATG

 14950 14960 14970 14980 14990 15000
 GTCAGATCCCTGGTCTGTTGCCGTCGAATCAGGCCAGTGTGATTGATGTTGATTCAT

 15010 15020 15030 15040 15050 15060
 ATATCGGTATTGATTCCAAGCGAAAGGCCAACGTGTTACCGTTACTTGATTGTGAAT

 15070 15080 15090 15100 15110 15120
 TAACTTCCCTTGAAGAGATGGCTTTGGTGGCGATACTTACGTTACGAGATCCACATTG

 15130 15140 15150 15160 15170 15180
 ATTCTGATGCACGTAACGGCGAGCAATTATTATTCTTCCATTACGATTGTTACGTAG

 15190 15200 15210 15220 15230 15240
 GGGATAAGAAGGTACTTATCATGCGTAATGGTGTGCTGGTTCTTACTGACGAAGAAC

 15250 15260 15270 15280 15290 15300
 TTTCTGATGGTAAAGCGTTATTCAACGACAAAGACAAAGCTGAGTTAGCAATGCTG

 15310 15320 15330 15340 15350 15360
 TTAAATCATCATTACGCCGTTATTACAACATAACCGTGGTCAATACGATTATAACGACA

 15370 15380 15390 15400 15410 15420
 TGATGAAGTTGGTTAATGGTATGTTGCCAGTTGTTGGTCCGAATATGATCAAGGTG

 15430 15440 15450 15460 15470 15480
 GCCGTAATCCATCATTGAAATTCTCGTCTGAGAAGTCTTGATGATTGAACGTATTACCA

 15490 15500 15510 15520 15530 15540
 AGATAGACCCAACCGGTGGTCATTGGGGACTAGGCCTGTTAGAAGGTCAAGAAAGATTAG

 15550 15560 15570 15580 15590 15600
 ACCCTGAGCATTGGTATTCCTTGTCACTTAAAGGTGATCAAGTAATGGCTGGTCGT

 15610 15620 15630 15640 15650 15660
 TGATGTCGGAAGGTTGTGGCAAATGGCGATGTTCTCATGCTGTCTGGTATGCATA

 15670 15680 15690 15700 15710 15720
 CCAATGTGAACAACGCTCGTTCCAACCACTACCAGGTGAATCACAAACGGTACGTTGTC

 15730 15740 15750 15760 15770 15780
 GTGGGCAAGTACTGCCACAGCGCAATACCTTAACCTACCGTATGGAAGTTACTGCGATGG

 15790 15800 15810 15820 15830 15840
 GTATGCATCCACAGCCATTGAAAGCTAATATTGATATTTGCTTGACGGTAAAGTGG

 15850 15860 15870 15880 15890 15900
 TTGTTGATTCAAAAACTTGAGCGTGATGATCAGCGAACAGATGAGCATTCAAGATTAC

 15910 15920 15930 15940 15950 15960
 CTGTAACACTGCCGAGTAATGTGGCGCTAAAGCGATTACTGCACCTGTTGCGTCAGTAG

Fig. 6

15970 15980 15990 16000 16010 16020
 CACCAGCATCTTCACCCGCTAACAGCGCGGATCTAGACGAACGTGGTGTGAACCGTTA

 16030 16040 16050 16060 16070 16080
 AGTTTCCTGAACGTCCGTTAATGCGTGGTGGACTCAGACTTGTCTGCACCGAAAAGCAAAG

 16090 16100 16110 16120 16130 16140
 GTGTGACACCGATTAAGCATTGAAAGCGCCTGCTGGTGTGGTCATCATAGAGTGCCTA

 16150 16160 16170 16180 16190 16200
 ACCAACGACCCGTTACACCTTGGCATATGTTGAGTTGCGACGGGTAATATTCTAACT

 16210 16220 16230 16240 16250 16260
 GTTTCGGTCCCTGATTTGATGTTATGAAGGTCGTATTCCACCTCGTACACCTTGTGGCG

 16270 16280 16290 16300 16310 16320
 ATTTACAAGTTGTTACTCAGGTTGAGAAGTGCAGGGCGAACGTCTTGATCTTAAAGATC

 16330 16340 16350 16360 16370 16380
 CATCAAGCTGTGTAGCTGAATACTATGTACCGGAAGACGCTTGGTACTTTACTAAAAACAA

 16390 16400 16410 16420 16430 16440
 GCCATGAAAATGGATGCCTTATTCAATTATGGAAATTGCATTGCAACCAAATGGCT

 16450 16460 16470 16480 16490 16500
 TTATTTCTGGTTACATGGGCACGACGCTTAAATACCCCTGAAAAAGATCTGTTCTCCGTA

 16510 16520 16530 16540 16550 16560
 ACCTTGATGGTAGCGGCACGTTATTAAAGCAGATTGATTTACCGGGCAAGACCATTGTGA

 16570 16580 16590 16600 16610 16620
 ATAAATCAGTCTGGTTAGTACGGCTATTGCTGGTGGCGCGATTATTCAAAGTTTACCGT

 16630 16640 16650 16660 16670 16680
 TTGATATGTCGTAGATGGCGAGCTATTATACTGGTAAAGCTGTATTGGTTACTTTA

 16690 16700 16710 16720 16730 16740
 GTGGTGAATCACTGACTAACCAACTGGCATTGATAACGGTAAACGACTAATGCGTGGT

 16750 16760 16770 16780 16790 16800
 TTGTTGATAACAATAACCCCGCAGCGAATATTGATGTGTTGATTAACTAATCAGTCAT

 16810 16820 16830 16840 16850 16860
 TGGCTCTGTATAAGCGCCTGCGATAAACCGCATTATAAATTGGCTGGTGGTCAGATGA

 16870 16880 16890 16900 16910 16920
 ACTTTATCGATACAGTGTCACTGGTGAAGGCAGGTGGTAAAGCAGGGCTGGCTATGTT

 16930 16940 16950 16960 16970 16980
 ATGGCGAACGTACGATTGATGCTGATGATTGGTCTTCCGTTATCACTTCCACCAAGATC

 16990 17000 17010 17020 17030 17040
 CGGTGATGCCAGGTTCAATTAGGTGTTGAAGCTATTATTGAGTTGATGCAGACCTATGCGC

 17050 17060 17070 17080 17090 17100
 TTAAAAATGATTGGTGGCAAGTTGCTAACCCACGTTCAATTGCGCCGATGACGCAAG

Fig. 6

17110 17120 17130 17140 17150 17160
 TTGATTGGAAATACCGTGGGCAAATTACGCCGCTGAATAAACAGATGTCAGTGGACGTGC

 17170 17180 17190 17200 17210 17220
 ATATCACTGAGATCGTGAATGACGCTGGTGAAGTGCAGATCGTGGTGAATCGAATCTGT

 17230 17240 17250 17260 17270 17280
 CTAAAGATGGCTCGCTATTATGAAGTTAAAACATCGTTAAGTATTGTTGAAGCGT

 17290 17300 17310 17320 17330 17340
 AAAGGGTCAAGTGTAAACGTGCTTAAGCGCCGCATTGGTAAAGACGCTTGCACGCCGTG

 17350 17360 17370 17380 17390 17400
 AATCCGTCATGGAGGCTGGGTTGGCATCCATGCCAACAAACAGCAAGCTTACTTTAAT

 17410 17420 17430 17440 17450 17460
 CAATACGGCTTGGTGTCCATTAGACGCCCGAACCTAGTTAATAGACAAAATAATT

 17470 17480 17490 17500 17510 17520
 TAGCTGTGGAATGAATATAGTAAGTAATCATTGGCAGCTACAAAAAAGGAATTAGAAT

 17530 17540 17550 17560 17570 17580
 GTCGAGTTAGGTTAACAAACAACGCAATTAACTGGGCTTGGAAAGTAGATCCAGC

 17590 17600 17610 17620 17630 17640
 GTCAGTTACACAAAGATGCAGAAATTAAAGCAGCTTAATGGATCTAACTAAACCTCT

 17650 17660 17670 17680 17690 17700
 CTATGTGGCGAACATAATTCAAGCGTAACGGTATAGCTAACATACGTCAGTAGCAGGTGC

 17710 17720 17730 17740 17750 17760
 GATCAGCAATAACATCGATGTTGATGTATTGGCGTTGCGAAAAGTTAAACCCAGAAGA

 17770 17780 17790 17800 17810 17820
 TCTGGGTGATGATGCTTACAAGAACAGCACGGCGTTAAATATGTTATCATGGCGGTGC

 17830 17840 17850 17860 17870 17880
 GATGGCAAATGGTATTGCCTCGGTTGAATTGGTTGCGTTAGGTAAAGCAGGGCTGTT

 17890 17900 17910 17920 17930 17940
 ATGTTCAATTGGTGCAGGTCTAGTGCCTGATGCGGTTGAAGATGCAATTGCGTAT

 17950 17960 17970 17980 17990 18000
 TCAAGCTGAATTACCAAATGCCCTTATGCCGTTAACTGATCCATGCACCCAGCAGAAGA

 18010 18020 18030 18040 18050 18060
 AGCATTAGAGCGTGGCGCGTTGAACGTTCTAAACTGGCGTCAAGACGGTAGAGGC

 18070 18080 18090 18100 18110 18120
 TTCAGCTTACCTGGTTAACTGAACACATTGTTGGTATCGTGCCTGGTCTAAACTAA

 18130 18140 18150 18160 18170 18180
 AACGCAGATGGCAGTGTAAATACGGTAACAAGGTTATCGCTAAAGTATCGCGTACCGA

 18190 18200 18210 18220 18230 18240
 AGTTGGTCGCCGCTTATGGAACCTGCACCGCAAAATTACTGGATAAGTTATTAGAAC

Fig. 6

18250 18260 18270 18280 18290 18300
 AAATAAGATCACCCCTGAACAAAGCTGCTTAGCGTTGCTTGTACCTATGGCTGATGATAT

 18310 18320 18330 18340 18350 18360
 TACTGGGAAAGCGGATTCTGGTGGTCATAACAGATAACCGTCCGTTAACATTATTACC

 18370 18380 18390 18400 18410 18420
 GACGATTATTGGTCTCGGTGATGAAGTGCAAGCGAAGTATAACTCTCCTGCATTACG

 18430 18440 18450 18460 18470 18480
 TGTTGGTGCTGGTGGTATCGAACGCCTGAAGCAGCACTCGCTGCATTAAACATGGG

 18490 18500 18510 18520 18530 18540
 CGCGGCTTATATCGTTCTGGGTTCTGTGAATCAGGCGTGTGTTGAAGCGGGTGCATCTGA

 18550 18560 18570 18580 18590 18600
 ATATACTCGTAAACTGTTATCGACAGTTGAAATGGCTGATGTGACTATGGCACCTGCTGC

 18610 18620 18630 18640 18650 18660
 AGATATGTTGAAATGGGTGTGAAGCTGCAAGTATTAAAACGCGGTCTATGTTCGCGAT

 18670 18680 18690 18700 18710 18720
 GCGTGCAGAAACTGTATGACTTGTATGTGGCTTATGACTCGATTGAAGATATCCCAGC

 18730 18740 18750 18760 18770 18780
 TGCTGAACGTGAGAAGATTGAAAAACAAATCTCCGTGAAACCTAGACGAGATTGGGA

 18790 18800 18810 18820 18830 18840
 TGGCACTATCGTTCTTACTGAACGCGATCCAGAAATGCTAGCCCGTGCACGAGTAG

 18850 18860 18870 18880 18890 18900
 TCCTAAACGTAAAATGGCACTTATCTCCGTTGGTATCTTGGCCTTCTCACGCTGGTC

 18910 18920 18930 18940 18950 18960
 AAACACAGGCGAGAAGGGACGTGAAATGGATTATCAGATTGGCAGGCCAAGTTAGG

 18970 18980 18990 19000 19010 19020
 TGCATTCAACAGCTGGGTGAAAGGTTCTTACCTTGAAGACTATAACCGCCGTGGCGCTGT

 19030 19040 19050 19060 19070 19080
 AGATGTTGCTTGCATATGCTTAAAGGTGCTGCGTATTTACAACGTGAAACCAGTTGAA

 19090 19100 19110 19120 19130 19140
 ATTGCAAGGTGTTAGCTTAAGTACAGAATTGGCAAGTTATCGTACGAGTGATTAATGTTA

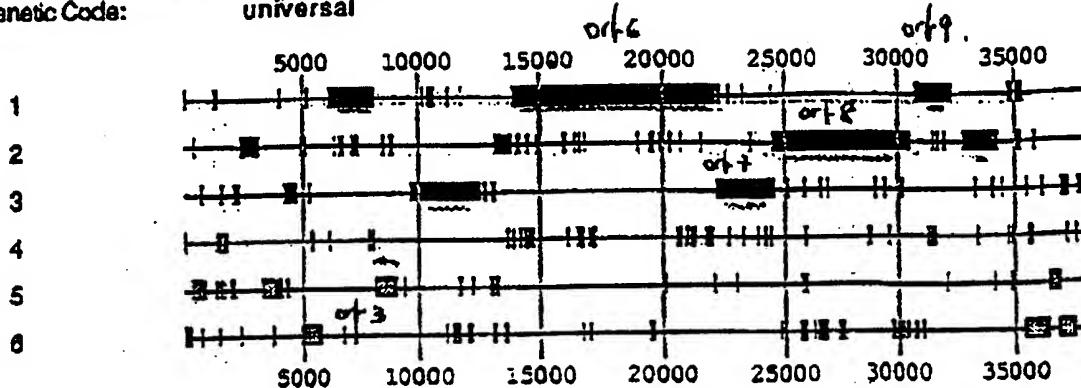
 19150 19160 19170 19180 19190 19200
 CTTGATGATATGTGAATTAATTAAAGCGCCTGAGGGCGCTTTTTGGTTTAACTCAG

 19210 19220
 GTGTTGTAACCGAAATTGCCCTTTC

Fig. 6

A

Start/Stop Method: AA span ≥ 25
 Genetic Code: universal



Page 1

B

Start/Stop Method: AA span ≥ 25
 Genetic Code: universal

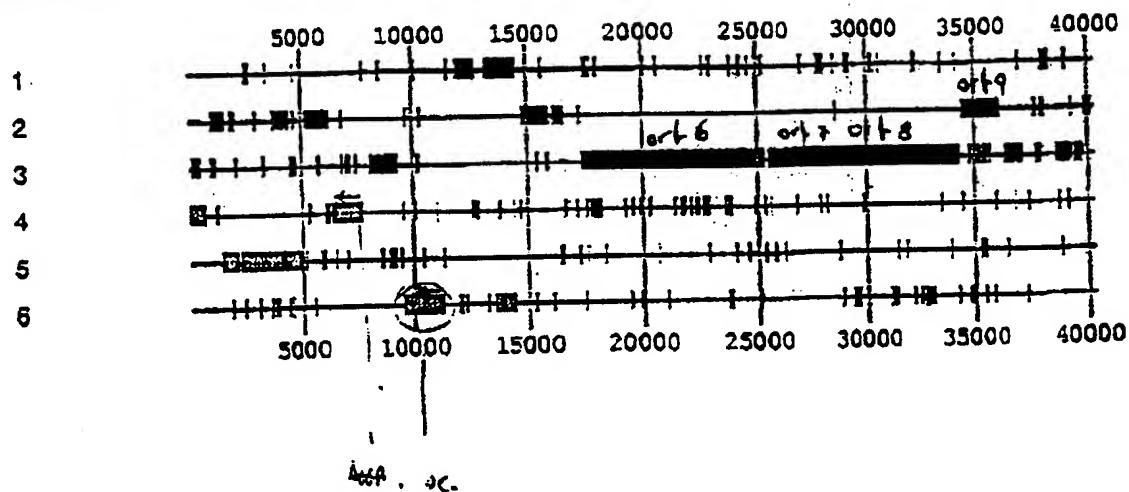


FIG 7

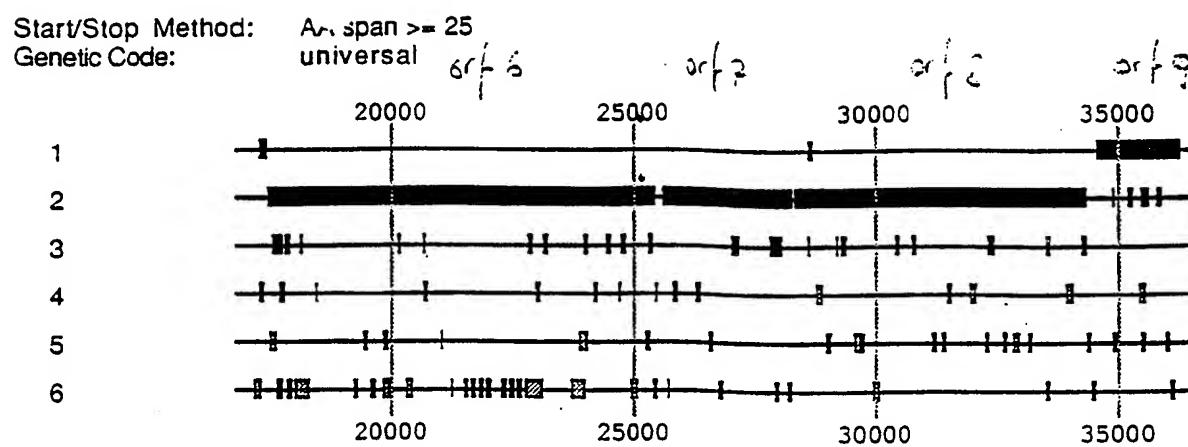


Fig. 8

Window Size = 8
Min. % Score = 60
Hash Value = 2

Scoring Matrix: BLOSUM 62

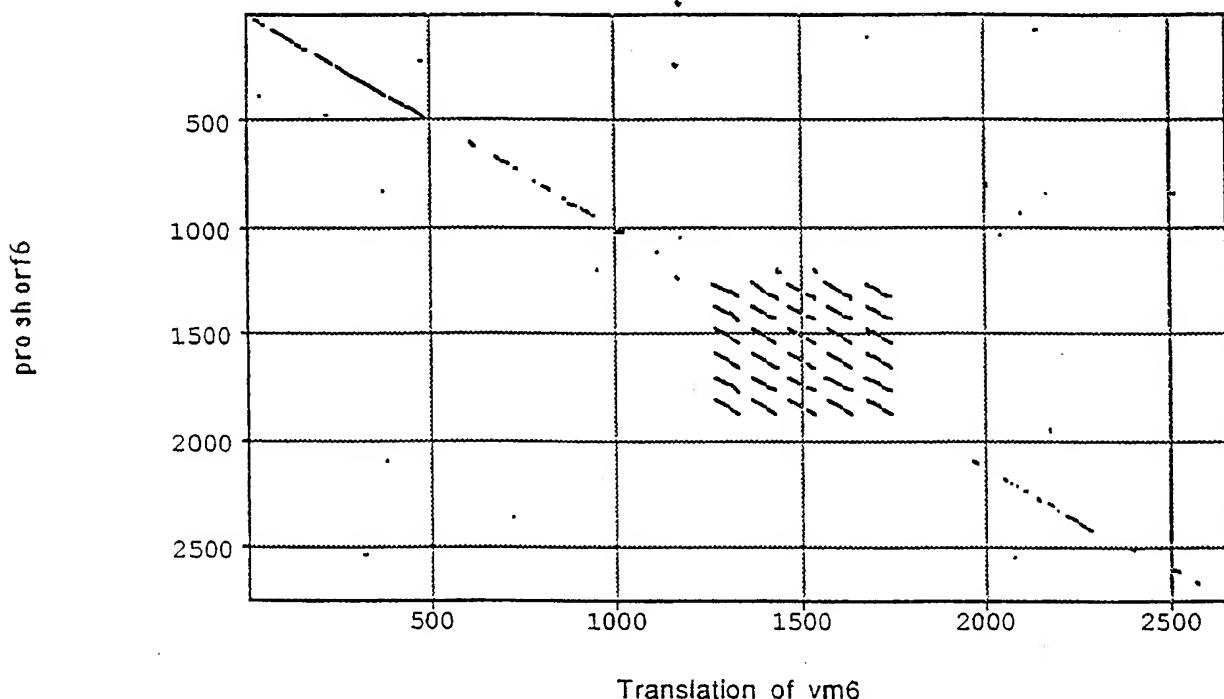


Fig. 9

Window Size = 8
Min. % Score = 60
Hash Value = 2

Scoring Matrix: BLOSUM 62

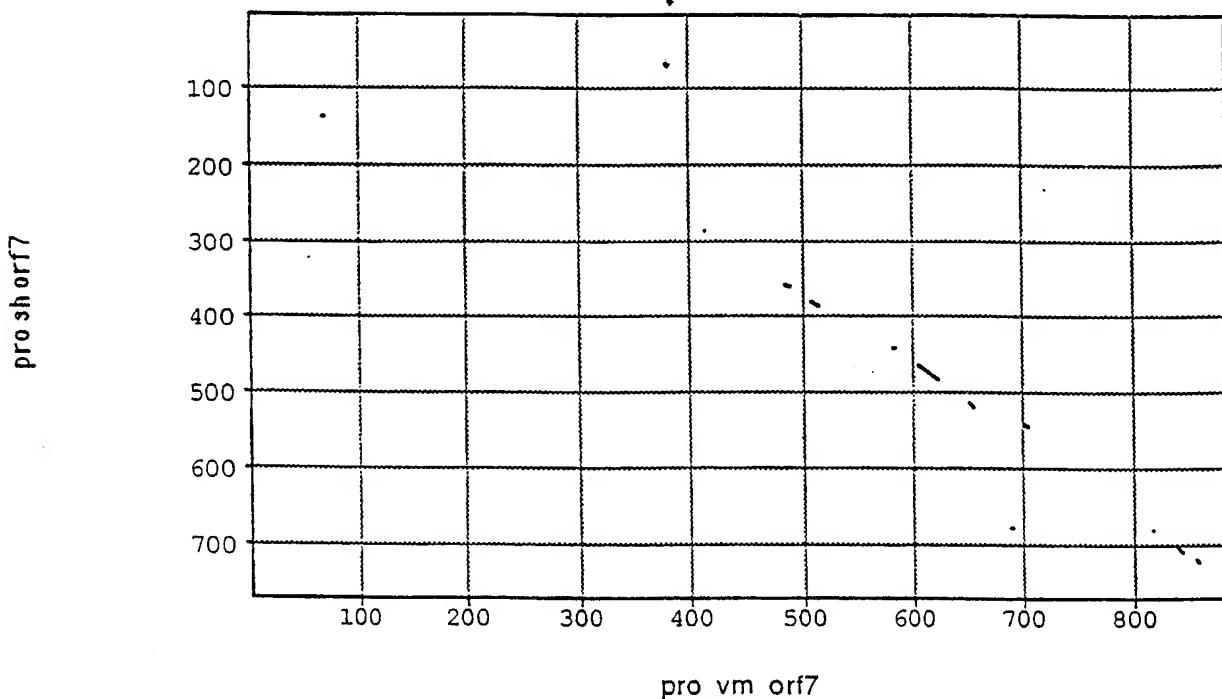


Fig. 10

Window Size = 8
Min. % Score = 60
Hash Value = 2

Scoring Matrix: BLOS 62

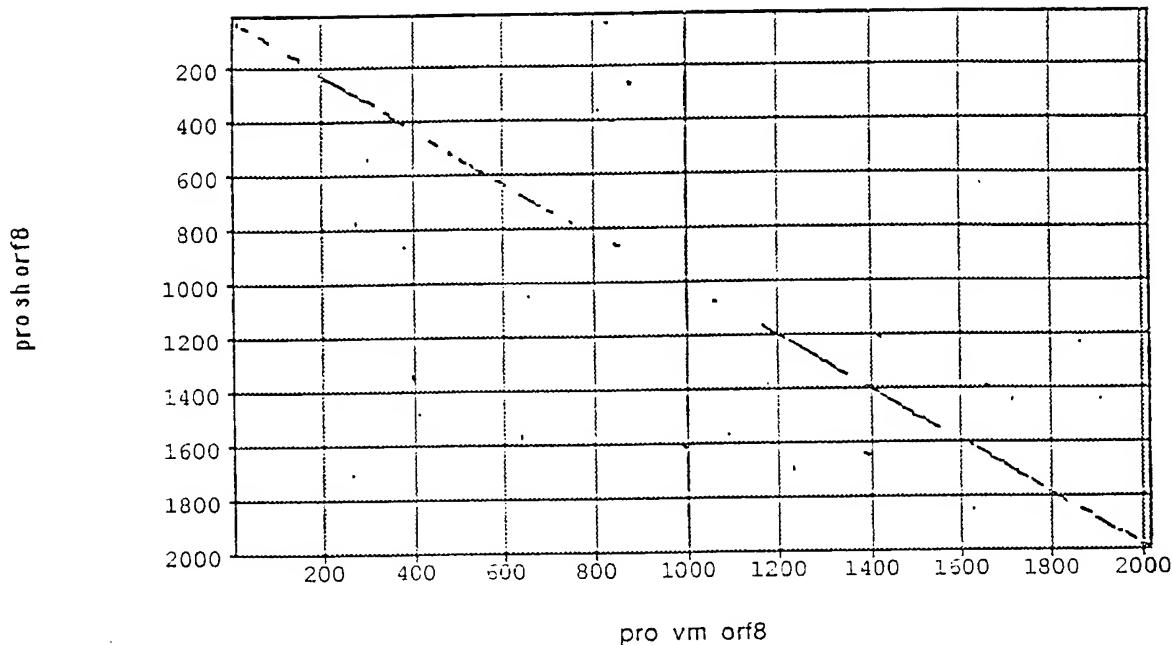


Fig. 11

Window Size = 8
Min. % Score = 60
Hash Value = 2

Scoring Matrix: BLOE . 62

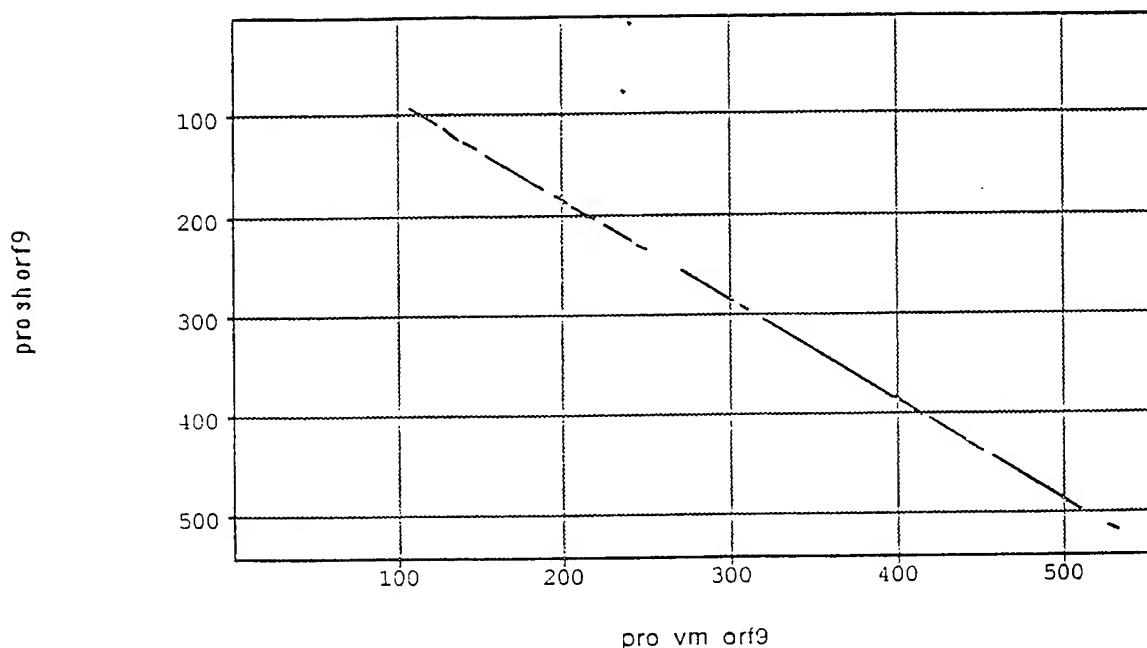


Fig. 12

COMPLEMENTATION Sp / Vm



Fig. 13

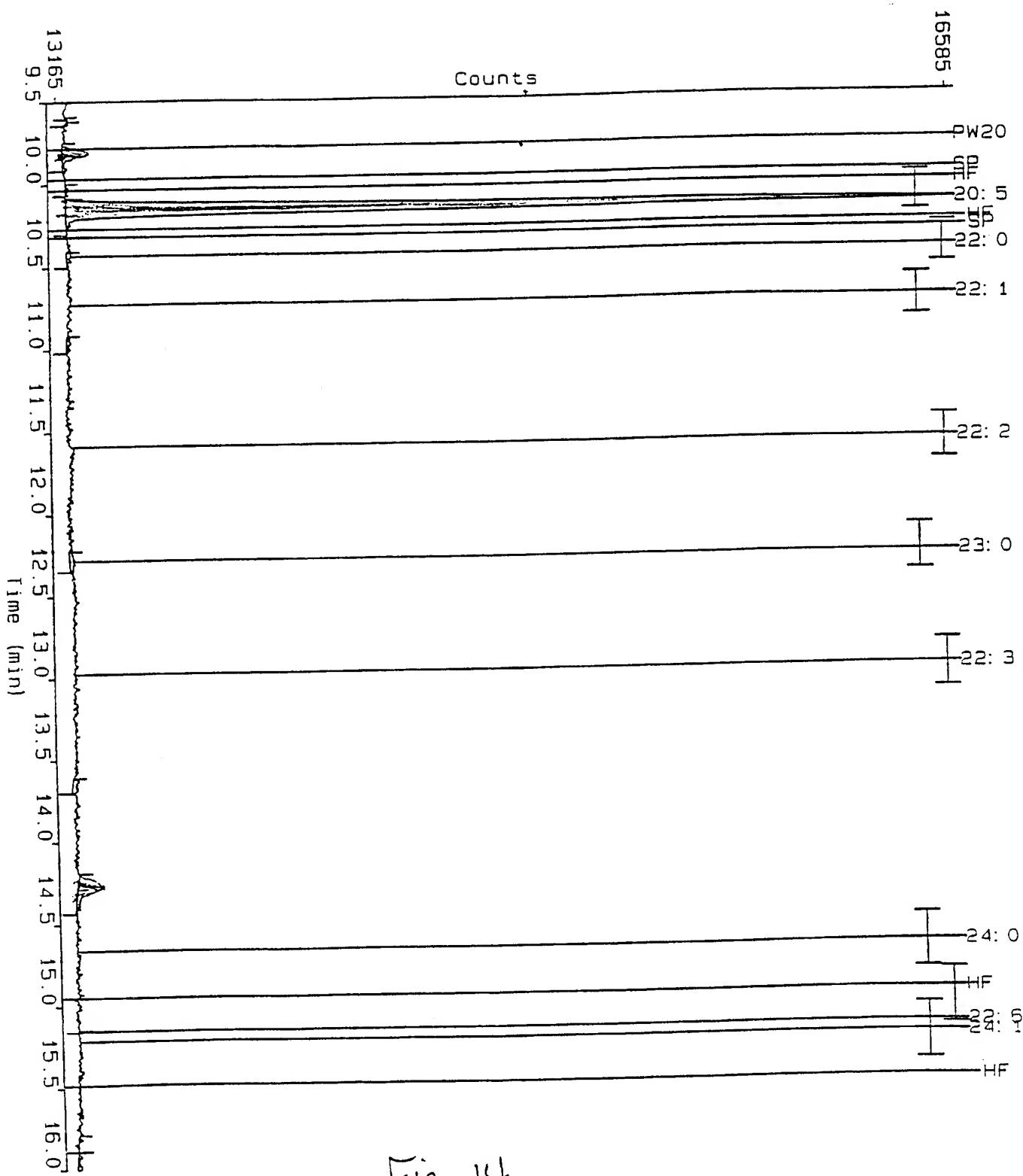


Fig. 14

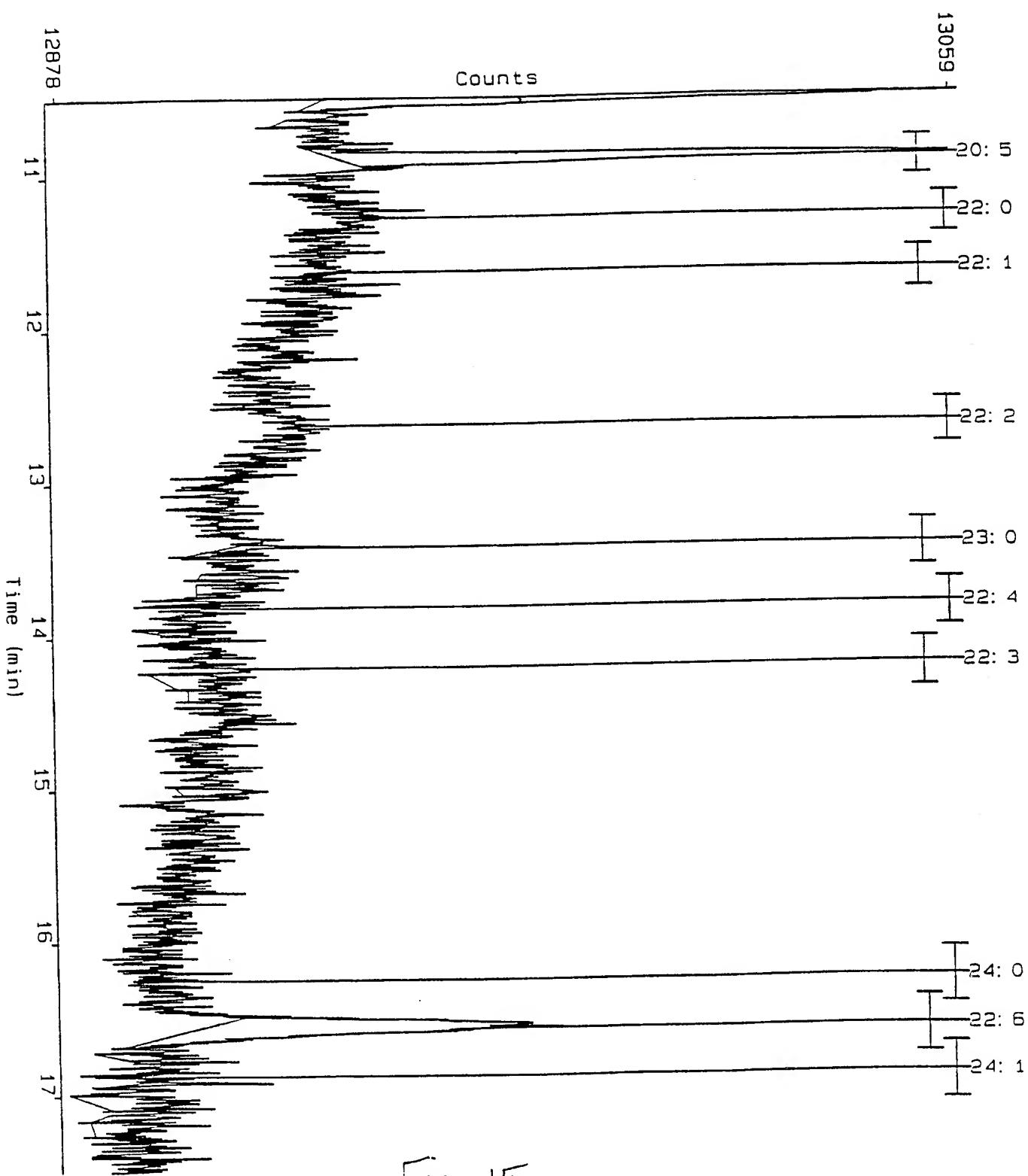


Fig. 15

<u>EPA (% Fatty acids)</u>	<u>DHA (% Fatty acids)</u>	<u>20°C</u>
0.00	0.06	pEPAD8
0.60	0.70	4
0.64	0.66	5
0.33	0.22	6s
0.45	0.59	6l
		<u>23°C</u>
0.02	0.06	pEPAD8
0.32	0.62	4
0.27	0.22	6s
0.18	0.65	6l

Fig. 16

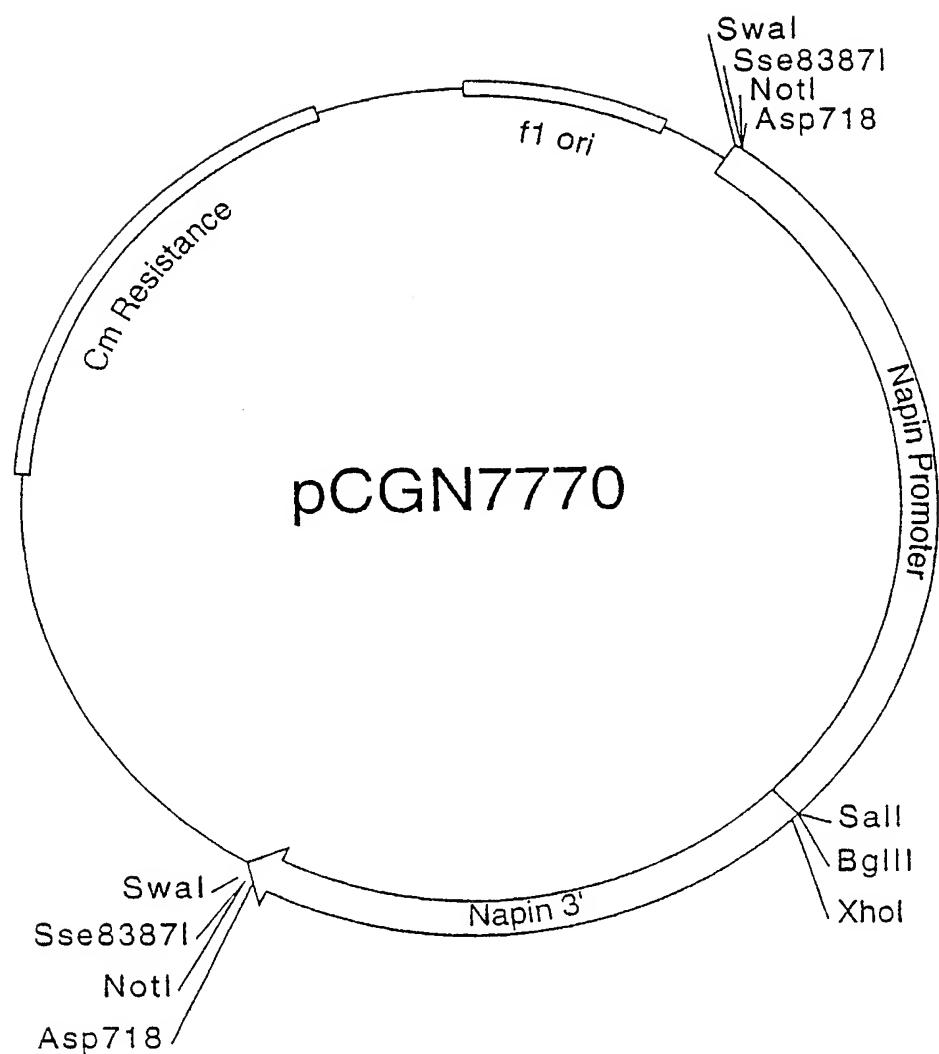


FIG 17

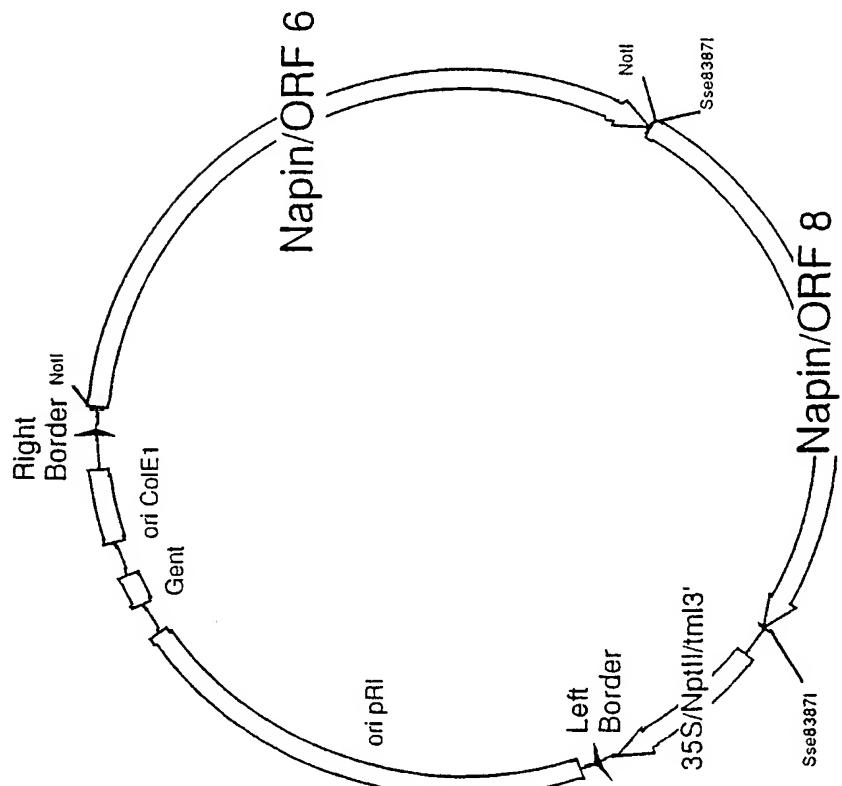
pCCG8535

FIG 18

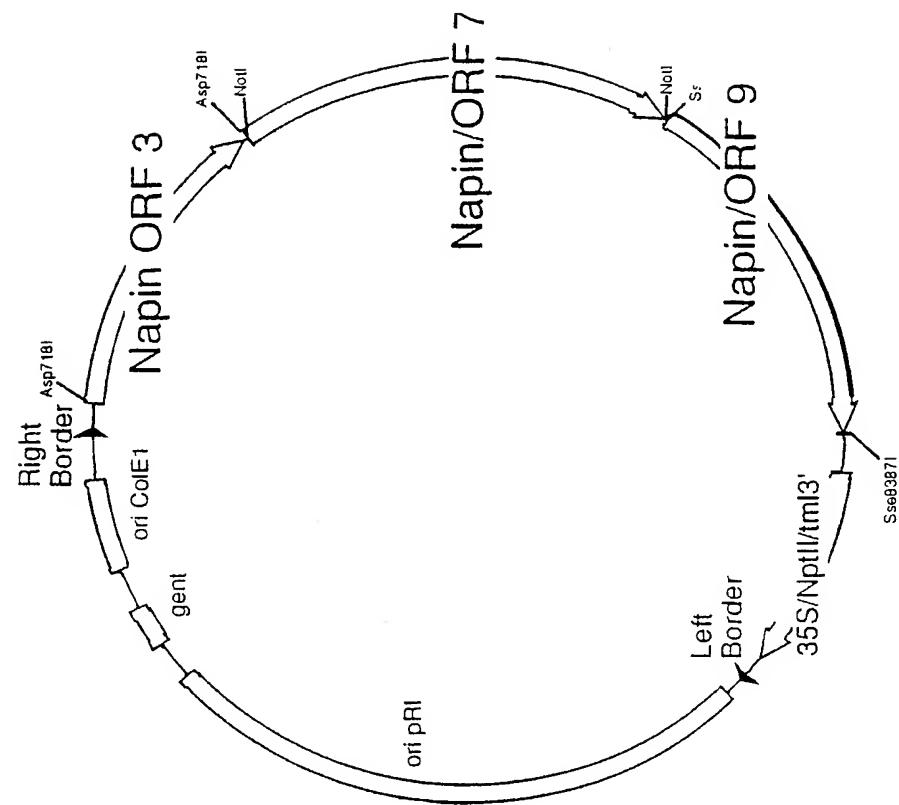
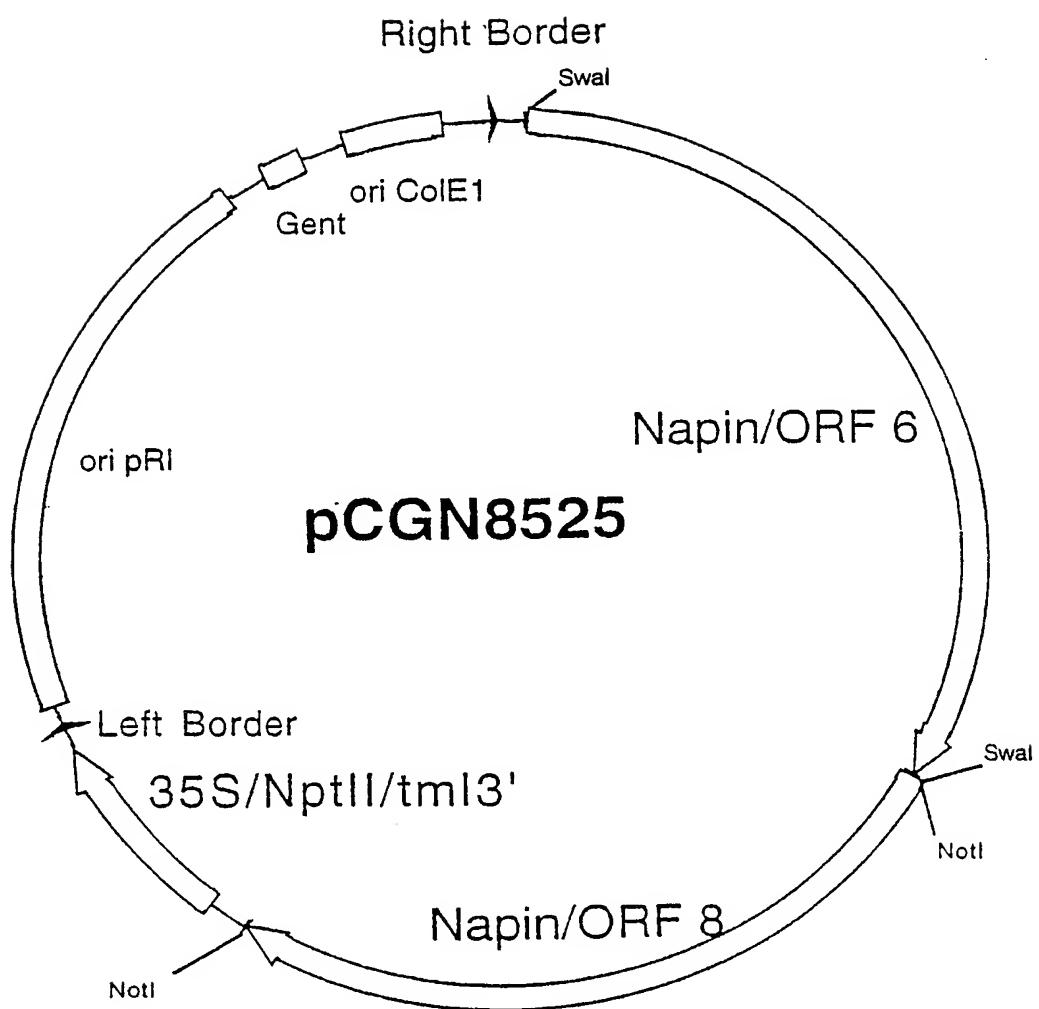
pCCG8537

FIG 19

**FIG 20**

YAZAWA(ORF1) (ORF2) (ORF3) (ORF4)(ORF5)(ORF6) (ORF7) (ORF8)

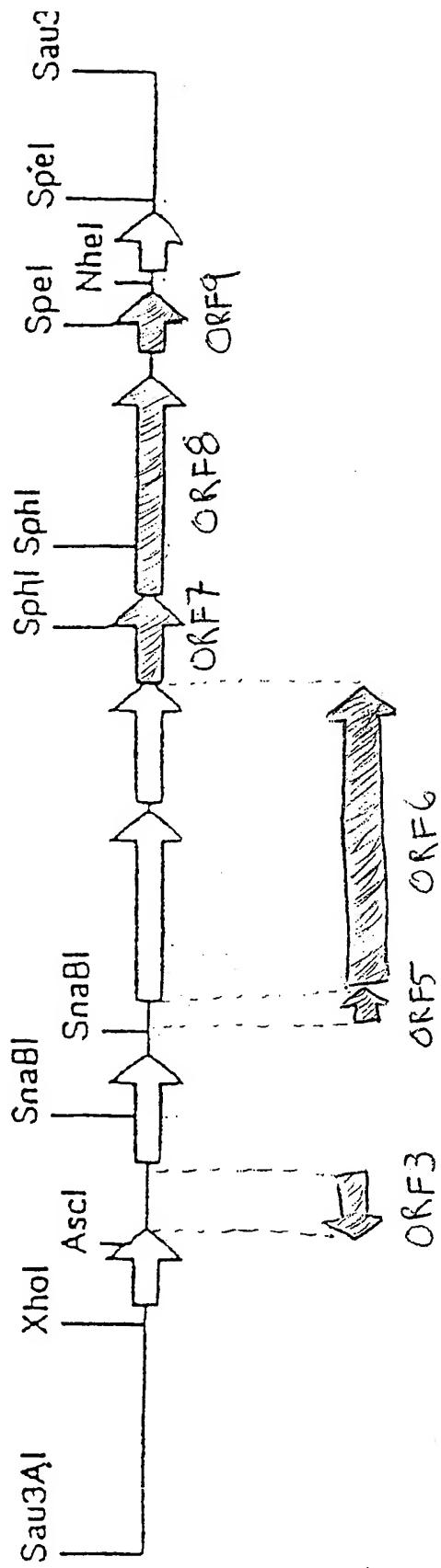


FIG 21

pCGN8556

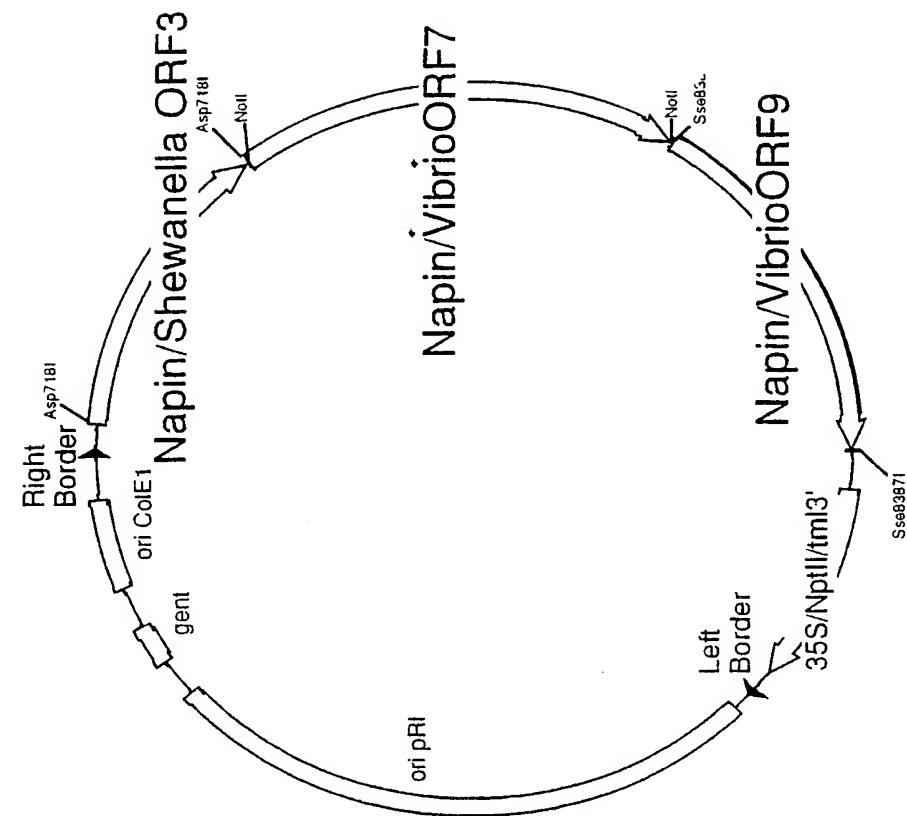


FIG 23

pCGN8560

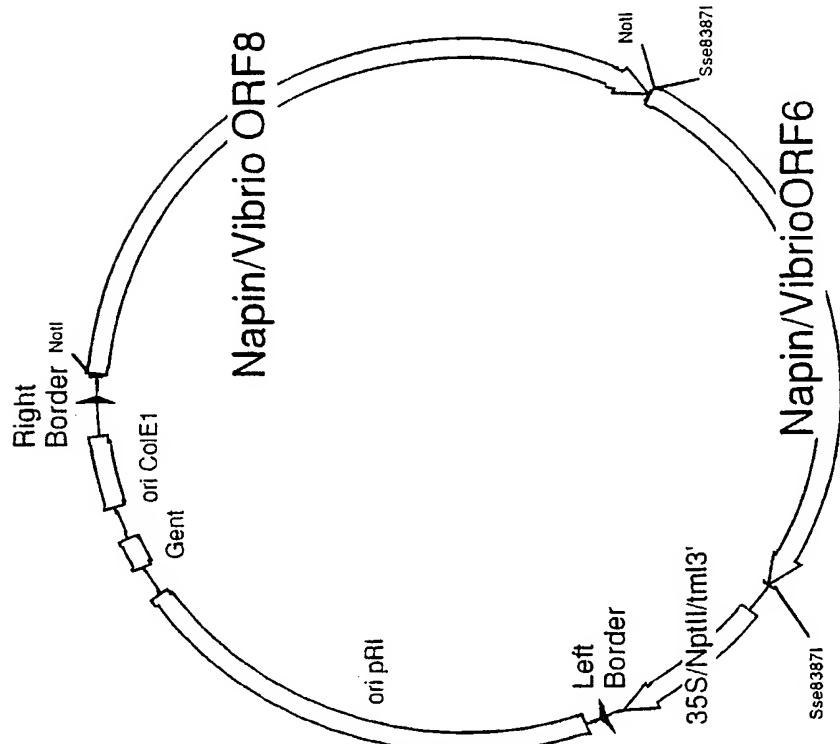


FIG 22

↓
ATT GGT AAA AAT AGG GGT TAT GTT TGT TGC TTT AAA GAG TGT CCT GAA
I G K N R G Y V C C F K E C P E>
↓ 9157 ↓
AAA TTG CTA ACT TCT CGA TTG ATT TCC TTA TAC TTC TGT CCG TTA ACA
K L L T S R L I S L Y F C P L T>
↓
ATA CAA GAG TGC GAT AAC CAG ACT ACA GAG TTG GTT AAG TCA TGG CTG
I Q E C D N Q T T E L V K S W L>
↓
CCT GAA GAT GAG TTA ATT AAG GTT AAT CGC TAC ATT AAA CAA GAA GCT
P E D E L I K V N R Y I K Q E A>
↓ 9016 ↓
AAA ACT CAA GGT TTA ATG GTA AGA G
K T Q G L M V R>

FIG 24

10	20	30	40	50	60
AGCGAAATGCTTATCAAGAAATTCCAAGATCAATACATCACTGGGAAGAAAATTCAATTCC					
70	80	90	100	110	120
CTGGTTCACTGGGTAACGTTATTCGGCGTATTGCTAACCGCTTCGACCTGGTGGCA					
130	140	150	160	170	180
TGAAC TGTCGTTGATGCAGCATGTGCAGGCCCTCTGCTCATGGTATGGCATTAA					
190	200	210	220	230	240
GCGAGCTTGTGAAGGCCGAGCGAAATGATGATTACAGGTGGTGTGTACCGATAACT					
250	260	270	280	290	300
CACCAACCATGTACATGAGCTCTCTAAACACCGGCATTACGACAAACGAAACAATT					
310	320	330	340	350	360
AACCATTGGATATTGACTCGAAAGGTATGATGATTGGTGAAGGTATCGGTATGATTGCGC					
370	380	390	400	410	420
TTAAACGTCTTGAAGACGCAGAGCGTGTGGCGACCGTATCTATTCCGTGATTAAAGGTG					
430	440	450	460	470	480
TTGGGTGCATCTCAGACGGTAATTATTAAGAGTANTTATGCGCNCCTGTCCTGAAGGTC					
490	500	510	520	530	540
AGGCTAAGGCACTTAACGTGCTTACGACGATGCAGGTTCGCACCGCACACACTGGCT					
550	560	570	580	590	600
TACTTGAAGGCCACGGCACAGGCACAGCAGCAGGTGATGTGGCAGAATTCACTGGTCCTA					
610	620	630	640	650	660
ACTCTGTATTCACTGAAAGGCAATGACGAAAAGCAACACATCGCATTAGGTTCACTGAAAT					
670	680	690	700	710	720
CACAGATTGGTCACACTAAATCAACAGCGGGTACTGCGGGTCAATCAAAGCGCTTTAG					
730	740	750	760	770	780
CACTGCACCAAAAGTACTGCCGCCAACATCAATGTAACCAGCCCTAACCTAAACTGA					
790	800	810	820	830	840
ATATTGAAGACTCGCCTTCTACCTCAATACACAGCGCTCCATGGATGCAACGTGTCG					
850	860	870	880		
ATGGTACACCGCGTCGTGGTATTAGCTATTGGTTGGT					

SS9 Photobacter

PCR Product Using Primers
Presented in Example I

FIG 25

Page 2

3-2(-VECTOR) by ORF, Insert Sequence
 Tuesday, November 24, 1998 11:06 PM

SEQUENCE RANGE: 4 50 600

10 20 30 40 50 60
 3-2(-VECTOR) CCACGTTAA GCACTTACG GTCCTATGA AGCTCCGTG TTTCCTGCGA AACAGCTGG
 1108 120 131 TO 509 OF ORF-2 A 1508 ... 160
 D. JNP11080 1030 1040 1050 1060 1070
 CCACGTTAA GCACTTACG GTCCTATGA AGCTCCGTG TTTCCTGCGA AACAGCTGG
 3-2(-VECTOR) CCACGTTAA GCACTTACG GTCCTATGA AGCTCCGTG TTTCCTGCGA AACAGCTGG

70 80 90 100 110 120
 3-2(-VECTOR) TCTAACTAA GCGCATGTA CGCTTACCGA AGCGCTTAA GCGCGCGATG TTTCCTGCG
 1708 180 191 TO 509 OF ORF-2 A 2108 220
 D. JNP11080 1090 1100
 TCTAACTAA GCGCATGTA C
 3-2(-VECTOR) TCTAACTAA GCGCATGTA C
 D. JNP1 st 1800 1810 1820
 GAG AGCTCCGTG TTTCCTGCGA AACAGCTGG
 3-2(-VECTOR) 1 1
 GAG AGCTCCGTG TTTCCTGCGA AACAGCTGG

130 140 150 160 170 180
 3-2(-VECTOR) GACCAACAG TTTCCTGCGA CGCTGCGAA ARAGCAGTGTG AGCGCGGTG GCTTCGTTA
 3308 340 351 TO 509 OF ORF-2 A 3708 ... 380
 D. JNP1 st 1180 1190
 G ATTCGCTGAG GTCGAGTAA
 3-2(-VECTOR)
 D. JNP1 st 1810
 GTCGAGTAA
 3-2(-VECTOR) GACCAACAG

190 200 210 220 230 240
 3-2(-VECTOR) ATCCGAAAT GGTCTACTA ATTCGCGCG TTGGTGTGTTG GGTATGTTA AGCGCGGT
 3908 400 411 TO 509 OF ORF-2 A 4108 ... 420
 D. JNP1 st 650 660
 G GGTATGTTA AGCGCGGT
 3-2(-VECTOR)
 D. JNP11080 1210 1220 1230
 ATCCGAAAT GGTCTACTA ATTCGCGCG AGCG
 3-2(-VECTOR) ATCCGAAAT GGTCTACTA ATTCGCGCG TTGG
 D. JNP1 st 2310 2320 2330 2340
 GGTCTACTA ATCCGAAAT GGTCTACTA ATTCGCGCG
 3-2(-VECTOR)
 GGTCTACTA ATCCGAAAT GGTCTACTA ATTCGCGCG

350 360 370 380 390 400
 3-2(-VECTOR) AGCGCGGT GGTCTACTA ATTCGCGCG GGTCTACTA GGTATGTTA ATTCGCGCG
 4508 460 471 TO 509 OF ORF-2 A 4708 ... 480
 D. JNP1 st 670
 AGCGCGGT
 3-2(-VECTOR) AGCGCGGT
 D. JNP1 st C
 3-2(-VECTOR) A
 D. JNP1 st 2900 2910 2920
 GGTCTACTA ATTCGCGCG GGTATGTTA ATTCGCGCG
 3-2(-VECTOR) 1 1
 GGTCTACTA ATTCGCGCG GGTATGTTA ATTCGCGCG

310 320 330 340 350 360
 3-2(-VECTOR) GGTATGTTA ATTCGCGCG GGTCTACTA ATTCGCGCG GGTCTACTA ATTCGCGCG
 4108 420 431 TO 509 OF ORF-2 A 4408 ... 450

ORF 6

Probe Resulting from PCR with Primers
 Presented in Example I

FIG 26A

3-1-VECT01 by ORF3 Aligned Sequences
Wednesday, November 25, 1998 11:06 PM

Page 3

0. Imp1 et	1340	1210	1300	1370
3-1-VECT01	GAGGCGG TTTTCTTAA CAGGAAAC CGCCGAGG TGTGCGCG			
	370	380	390	400
3-1-VECT01	AGAGGTTG CGCGGTTG CGGGTTTG CGCTTGGG TTGGCGCG			
	4784	101	101 TO 509	OF ORF3-2
	3084			
0. Imp1&2	1398	1400	1410	1420
3-1-VECT01	AGAGGTTG CGCGGTTG CGGGTTTG CGCTTGGG TTGGCGCG			

FIG 26B

INTERNATIONAL SEARCH REPORT

Int	International Application No
PCT/US 98/11639	

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C12N15/31 C12N15/52 C12N15/82 C12N15/70 C12N5/10
 C12N1/21 C12P7/64 A01H5/00

According to International Patent Classification(IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
 IPC 6 C12N C12P C07K A01H

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	NAKAHARA, TORO: "Physiological activity of docosahexaenoic acid (DHA) and its production by microbial culture" YUKAGAKU (1995), 44(10), 821-7 CODEN: YKGKAM; ISSN: 0513-398X, XP002080682 see abstract ---	6,7, 11-13
A		14,32
X	NASU M ET AL: "Efficient transformation of Marchantia polymorpha that is haploid and has very small genome DNA; Agrobacterium tumefaciens-mediated transformation of suspension cell culture, for use in eicosapentaenoic acid, arachidonic acid and antibiotic production" J.FERMENT.BIOENG.;(1997) 84, 6, 519-23 CODEN: JFBIEX ISSN: 0922-338X, XP002080470 see the whole document ---	25,27, 28,30
		-/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

° Special categories of cited documents :

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- "O" document referring to an oral disclosure, use, exhibition or other means
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"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

Date of mailing of the international search report

14 October 1998

23/10/1998

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Authorized officer

Kania, T

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 98/11639

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	KYLE D ET AL: "Long-chain omega-3 polyunsaturated fatty acids: prospects for introduction into horticultural food plants; e.g. alga eicosapentaenoic acid and docosahexaenoic acid gene cloning, expression in transgenic plant oil, crop improvement (conference paper)" HORTSCIENCE;(1990) 25, 12, 1523-26 CODEN: HJHSAR, XP002080471 * see the whole document, esp. p.1524, 2nd par. * ---	25-28, 30,31
X	EP 0 594 868 A (SAGAMI CHEM RES) 4 May 1994 cited in the application see the whole document ---	15-17, 19-22,24
X	WO 96 21735 A (SAGAMI CHEM RES) 18 July 1996 cited in the application see the whole document ---	15-17, 19-22,24
A	YAZAWA, KAZUNAGA: "Production of eicosapentaenoic acid from marine bacteria" LIPIDS (1996), 31(SUPPL., FATTY ACIDS AND LIPIDS FROM CELL BIOLOGY TO HUMAN DISEASE), S297-S300 CODEN: LPDSAP;ISSN: 0024-4201, XP002080483 cited in the application see the whole document ---	1-32
A	SOMERVILLE C R: "Future prospects for genetic modification of the composition of edible oils from higher plants; oilseed crop improvement by lipid and fatty acid modification (conference paper)" AM.J.CLIN.NUTR.;(1993) 58, 2, SUPPL., 270S-275S CODEN: AJCNAC, XP002080472 * see esp. p.274S, r. col., 1st par. * -----	1-32

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 98/11639

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